

\*\*\*\*\* RECONNECTED TO STN INTERNATIONAL \*\*\*\*\*  
 SESSION RESUMED IN FILE 'PHAR' AT 15:38:09 ON 24 JAN 2003  
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 COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE	TOTAL
ENTRY	SESSION
40.86	265.93

=> index bioscience medicine  
 FILE 'DRUGMONOG' ACCESS NOT AUTHORIZED  
 COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE	TOTAL
ENTRY	SESSION
44.02	269.09

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DRUGB, DRUGLAUNCH, DRUGMONOG2, ...' ENTERED AT 15:38:32 ON 24 JAN 2003

67 FILES IN THE FILE LIST IN STNINDEX

Enter SET DETAIL ON to see search term postings or to view  
 search error messages that display as 0\* with SET DETAIL OFF.

=> s exendin and glucagon?

44 FILE ADISCTI  
 7 FILE ADISINSIGHT  
 3 FILE ADISNEWS  
 6 FILE AGRICOLA  
 2 FILE AQUASCI  
 2 FILE BIOBUSINESS  
 2 FILE BIOCOMMERCE  
 222 FILE BIOSIS  
 0\* FILE BIOTECHABS

10 FILES SEARCHED...

3 FILE BIOTECHDS  
 53 FILE BIOTECHNO  
 15 FILE CABA  
 27 FILE CANCERLIT  
 194 FILE CAPLUS  
 22 FILE CIN  
 0\* FILE DDFU

23 FILES SEARCHED...

1185 FILE DGENE  
 4 FILE DRUGNL  
 116 FILE DRUGU  
 4 FILE DRUGUPDATES  
 5 FILE EMBAL  
 147 FILE EMBASE  
 117 FILE ESBIODASE  
 2 FILE FEDRIP  
 2 FILE FROSTI  
 2 FILE GENBANK  
 8 FILE IFIPAT  
 6 FILE JICST-EPLUS  
 24 FILE LIFESCI  
 156 FILE MEDLINE

48 FILES SEARCHED...

74 FILE PASCAL  
 3 FILE PHAR  
 4 FILE PHARMAML  
 12 FILE PHIN  
 31 FILE PROMT  
 181 FILE SCISEARCH  
 55 FILE TOXCENTER

62 FILE USPATFULL  
4 FILE USPAT2  
25 FILE WPIDS  
0\* FILE WPINDEX  
64 FILES SEARCHED...  
1 FILE IPA  
18 FILE NLDB

40 FILES HAVE ONE OR MORE ANSWERS, 67 FILES SEARCHED IN STNINDEX

L309 QUE EXENDIN AND GLUCAGON?

=> file hits

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
31.35	300.44

FULL ESTIMATED COST

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FILE 'IPA' ENTERED AT 16:12:37 ON 24 JAN 2003

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=> s 1309

L310	1185	FILE	DGENE
L311	222	FILE	BIOSIS
L312	194	FILE	CAPLUS
L313	181	FILE	SCISEARCH
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L320	55	FILE	TOXCENTER
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L346	2	FILE	FEDRIP
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L348	2	FILE	GENBANK
L349	1	FILE	IPA

TOTAL FOR ALL FILES

L350 2850 L309

=> s exendin (s) glucagon

L351	935	FILE	DGENE
L352	176	FILE	BIOSIS
L353	105	FILE	CAPLUS
L354	158	FILE	SCISEARCH
L355	139	FILE	MEDLINE
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L359	67	FILE	PASCAL
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L383	1	FILE	PHAR
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L385	0	FILE	BIOBUSINESS
L386	2	FILE	BIOCOMMERCE
PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH			
FIELD CODE - 'AND' OPERATOR ASSUMED 'EXENDIN (S) GLUCAGON'			
L387	2	FILE	FEDRIP
L388	2	FILE	FROSTI
L389	0	FILE	GENBANK
L390	1	FILE	IPA

TOTAL FOR ALL FILES

L391	2223	EXENDIN (S) GLUCAGON
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=> s L391 and exendin-4 and glucagon

L392	396	FILE	DGENE
L393	103	FILE	BIOSIS
L394	77	FILE	CAPLUS
L395	109	FILE	SCISEARCH
L396	84	FILE	MEDLINE
L397	75	FILE	EMBASE
L398	65	FILE	ESBIOBASE
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L418	3	FILE DRUGNL
L419	3	FILE DRUGUPDATES
L420	4	FILE PHARMAML
L421	2	FILE USPAT2
L422	2	FILE ADISNEWS
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L424	1	FILE PHAR
L425	1	FILE AQUASCI
L426	0	FILE BIOBUSINESS
L427	1	FILE BIOCOMMERCE
L428	2	FILE FEDRIP
L429	1	FILE FROSTI
L430	0	FILE GENBANK
L431	1	FILE IPA

TOTAL FOR ALL FILES

L432 1243 L391 AND EXENDIN-4 AND GLUCAGON

=> s l432 and glucagonoma

L433	0	FILE DGENE
L434	0	FILE BIOSIS
L435	1	FILE CAPLUS
L436	0	FILE SCISEARCH
L437	0	FILE MEDLINE
L438	0	FILE EMBASE
L439	0	FILE ESBIODBASE
L440	0	FILE DRUGU
L441	0	FILE PASCAL
L442	1	FILE USPATFULL
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L459	0	FILE DRUGNL
L460	0	FILE DRUGUPDATES
L461	0	FILE PHARMAML
L462	0	FILE USPAT2
L463	0	FILE ADISNEWS
L464	0	FILE BIOTECHDS
L465	0	FILE PHAR
L466	0	FILE AQUASCI
L467	0	FILE BIOBUSINESS
L468	0	FILE BIOCOMMERCE
L469	0	FILE FEDRIP
L470	0	FILE FROSTI
L471	0	FILE GENBANK
L472	0	FILE IPA

TOTAL FOR ALL FILES

L473 5 L432 AND GLUCAGONOMA

=> d l473 1-4 ibib abs

L473 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2003 ACS  
 ACCESSION NUMBER: 2000:493318 CAPLUS  
 DOCUMENT NUMBER: 133:129880  
 TITLE: Methods using an **exendin** or related substance for **glucagon** suppression  
 INVENTOR(S): Young, Andrew; Gedulin, Bronislava  
 PATENT ASSIGNEE(S): Amylin Pharmaceuticals, Inc., USA  
 SOURCE: PCT Int. Appl., 96 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 3  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000041548	A2	20000720	WO 2000-US942	20000114
WO 2000041548	A3	20001130		
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1143989	A2	20011017	EP 2000-902415	20000114
EP 1143989	A3	20020911		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
BR 2000007823	A	20011120	BR 2000-7823	20000114
JP 2002538084	T2	20021112	JP 2000-593169	20000114
NO 2001003469	A	20010914	NO 2001-3469	20010712
PRIORITY APPLN. INFO.:				
			US 1999-116380P	P 19990114
			US 1999-132017P	P 19990430
			US 2000-175365P	P 20000110
			WO 2000-US942	W 20000114

AB Methods are provided for use of an **exendin**, an **exendin** agonist, or a modified **exendin** or **exendin** agonist having an **exendin** or **exendin** agonist linked to one or more polyethylene glycol polymers, for example, for lowering **glucagon** levels and/or suppressing **glucagon** secretion in a subject. These methods are useful in treating hyperglucagonemia and other conditions that would be benefited by lowering plasma **glucagon** or suppressing **glucagon** secretion.

L473 ANSWER 2 OF 5 USPATFULL  
 ACCESSION NUMBER: 2003:4123 USPATFULL  
 TITLE: Use of glycogen phosphorylase inhibitors  
 INVENTOR(S): Treadway, Judith L., Mystic, CT, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003004162	A1	20030102
APPLICATION INFO.:	US 2001-813335	A1	20010320 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-191381P	20000322 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Gregg C. Benson, Pfizer Inc., Patent Department, MS 4159,, Eastern Point Road, Groton, CT, 06340	
NUMBER OF CLAIMS:	23	

EXEMPLARY CLAIM: 1  
LINE COUNT: 4011  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides methods of treating prophylactically an individual in whom Type 2 diabetes mellitus has not yet presented, but in whom there is an increased risk of developing such condition, which methods comprise administering to an individual in need thereof an effective amount of a glycogen phosphorylase inhibitor; effective amounts of a glycogen phosphorylase inhibitor and a non-glycogen phosphorylase inhibiting anti-diabetic agent; or effective amounts of a glycogen phosphorylase inhibitor and an anti-obesity agent.

The invention further provides methods of treating prophylactically an individual in whom Type 2 diabetes mellitus has not yet presented, but in whom there is an increased risk of developing such condition, which methods comprise administering to an individual in need thereof a pharmaceutical composition comprising effective amounts of a glycogen phosphorylase inhibitor and a non-glycogen phosphorylase inhibiting anti-diabetic agent; or effective amounts of a glycogen phosphorylase inhibitor and an anti-obesity agent.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L473 ANSWER 3 OF 5 WPIDS (C) 2003 THOMSON DERWENT

ACCESSION NUMBER: 2002-012518 [02] WPIDS  
CROSS REFERENCE: 2000-595483 [50]; 2000-680964 [50]  
DOC. NO. CPI: C2002-003289  
TITLE: Use of glycogen phosphorylase inhibitor in prophylactic treatment of Type II diabetes.  
DERWENT CLASS: B02  
INVENTOR(S): TREADWAY, J L  
PATENT ASSIGNEE(S): (PFIZ) PFIZER PROD INC; (TREA-I) TREADWAY J L  
COUNTRY COUNT: 34  
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
EP 1136071	A2	20010926	(200202)*	EN	78
R: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT LU LV MC MK NL PT					
RO SE SI TR					
AU 2001028130	A	20010927	(200202)		
CA 2341344	A1	20010922	(200203)	EN	
JP 2001302546	A	20011031	(200204)		70
HU 2001001158	A2	20020228	(200223)		
KR 2001092696	A	20011026	(200223)		
NZ 510677	A	20021025	(200274)		
US 2003004162	A1	20030102	(200305)		
ZA 2001002318	A	20021127	(200305)		154

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
EP 1136071	A2	EP 2001-301979	20010305
AU 2001028130	A	AU 2001-28130	20010320
CA 2341344	A1	CA 2001-2341344	20010320
JP 2001302546	A	JP 2001-78839	20010319
HU 2001001158	A2	HU 2001-1158	20010321
KR 2001092696	A	KR 2001-14306	20010320
NZ 510677	A	NZ 2001-510677	20010321
US 2003004162	A1 Provisional	US 2000-191381P	20000322
		US 2001-813335	20010320
ZA 2001002318	A	ZA 2001-2318	20010320



PRIORITY APPLN. INFO: US 2000-191381P 20000322; US 2001-813335  
20010320

AN 2002-012518 [02] WPIDS  
CR 2000-595483 [50]; 2000-680964 [50]  
AB EP 1136071 A UPAB: 20020114

NOVELTY - A glycogen phosphorylase inhibitor (G1) is used in the manufacture of a medicament for prophylactically treating an individual with increased risk of developing Type II diabetes mellitus  
DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the following:

- (1) a pharmaceutical composition comprising (G1) and a non-glycogen phosphorylase inhibiting anti-diabetic agent (NG1); and
- (2) a pharmaceutical composition comprising (G1) and an anti-obesity agent.

ACTIVITY - Antidiabetic.

MECHANISM OF ACTION - Glycogen phosphorylase inhibitor.

No biological data is given.

USE - For prophylactically treating a person having risk associated with Type 2 diabetes (particularly risk associated with insulin resistance and/or hyperinsulinemia; environmental or genetic Type 2 diabetes predisposing disease states or conditions (e.g. person with a family history of diabetes); race and/or ethnicity (e.g. individuals from African-American, Hispanic, Native American, Asian, or Pacific Islander population); genetic mutations affecting beta -cell function (e.g. defect on chromosome 12, gene HNF-1 alpha (MODY3), chromosome 7, gene glucokinase (MODY2), chromosome 20, gene HNF-4a (MODY1), or mitochondrial DNA); genetic defects in insulin action (e.g. genetic mutation leading to Type A insulin resistance, acanthosis nigricans, leprechaunism, Rabson-Mendenhall syndrome; lipotrophic diabetes, or a genetic mutation or mutations in the insulin receptor, IRS proteins, glucose transporters, PC-1, glucokinase, UCP-1, beta 3 adrenergic receptor gene); presence of excess adipose tissue or clinically diagnosed obesity (e.g. central obesity); clinical chemistry or diagnostic testing signifying a pre-diabetic state (e.g. impaired glucose tolerance, impaired fasting glucose, or hyperglycemia relative to normoglycemia); physiologic and endocrine changes associated with growth, development, or aging (e.g. menopausal, pubescent, or aged individuals); diet or eating behaviors (e.g. consumption of high fat or high carbohydrate diets, experiencing prolonged fasting or starvation, having anorexia nervosa and bulimia); abnormal cardiovascular or blood lipid parameters (e.g. hypertension, HDL cholesterol level upto 35 mg/dl and/or TG levels of at least 250 mg/dl and metabolic syndrome); reproductive status (e.g. pregnancy, a history of gestational diabetes and macrosomia); muscle wasting (e.g. aging, starvation, exposure to anti-gravity environments and paralysis resulting from spinal cord injury); polycystic ovary syndrome; organ disease or dysfunction (e.g. liver cirrhosis and renal disease); metabolic disturbances; endocrine disorders or endocrinopathies (e.g. hyperandrogenism, thyrotoxicosis, hyperthyroidism, insulinoma, **glucagonoma**, somatostatinoma, aldosteroma, Cushing's Syndrome, pheochromocytoma, acromegaly and hypercortisolemia); pathophysiologic states (e.g. infection, congenital rubella, cytomegalovirus, toxemia, uremia, sepsis and trauma); immune-mediated disease (e.g. stiff man syndrome or the production of anti-insulin receptor antibodies); drug or chemical exposure (e.g. glucocorticoids, cytokines, alpha -interferon, thyroid hormone, TNF alpha , thiazides, estrogen-containing products, beta -blockers, nicotinic acid, serotonin receptor-targeted antipsychotics or antidepressants, vacor, diazoxide, dilantin, and HIV protease inhibitors); genetic syndrome associated with diabetes (e.g. Down's Syndrome, Klinefelter's Syndrome, Wolfram's Syndrome, Freidreich's Syndrome, Huntington's chorea, Laurence-Moon-Biedl Syndrome, myotonic dystrophy, porphyria, Prader-Willi Syndrome and Alzheimer's Disease); and detrimental effects caused by the administration of prolonged, elevated doses of insulin and/or the presence of ketoacidosis) (all claimed).  
Dwg.0/0

ACCESSION NUMBER: 2000-490999 [43] WPIDS  
 CROSS REFERENCE: 2000-514584 [46]; 2001-514422 [56]  
 DOC. NO. CPI: C2000-147547  
 TITLE: Lowering plasma glucagon using exendin  
 , an exendin agonist, a modified  
 exendin or a modified exendin agonist,  
 usefull for treating hyperglucagonemia and diabetes.  
 DERWENT CLASS: A25 A96 B04  
 INVENTOR(S): GEDULIN, B; YOUNG, A  
 PATENT ASSIGNEE(S): (AMYL-N) AMYLIN PHARM INC  
 COUNTRY COUNT: 91  
 PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 2000041548	A2	20000720	(200043)*	EN	96
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FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS					
LT LU LV MA MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL					
TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW					
AU 2000024136	A	20000801	(200054)		
NO 2001003469	A	20010914	(200163)		
EP 1143989	A2	20011017	(200169)	EN	
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RO SE SI					
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KR 2001086165	A	20010908	(200219)		
KR 2002001719	A	20020109	(200246)		
CN 1347327	A	20020501	(200252)		
JP 2002538084	W	20021112	(200275)		104

#### APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 2000041548	A2	WO 2000-US942	20000114
AU 2000024136	A	AU 2000-24136	20000114
NO 2001003469	A	WO 2000-US942	20000114
		NO 2001-3469	20010712
EP 1143989	A2	EP 2000-902415	20000114
		WO 2000-US942	20000114
BR 2000007823	A	BR 2000-7823	20000114
		WO 2000-US942	20000114
KR 2001086165	A	KR 2001-708904	20010713
KR 2002001719	A	WO 2000-US942	20000114
		KR 2001-708892	20010713
CN 1347327	A	CN 2000-805017	20000114
JP 2002538084	W	JP 2000-593169	20000114
		WO 2000-US942	20000114

#### FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 2000024136	A Based on	WO 200041548
EP 1143989	A2 Based on	WO 200041548
BR 2000007823	A Based on	WO 200041548
KR 2002001719	A Based on	WO 200041548
JP 2002538084	W Based on	WO 200041548

PRIORITY APPLN. INFO: US 2000-175365P 20000110; US 1999-116380P  
 19990114; US 1999-132017P 19990430  
 AN 2000-490999 [43] WPIDS

CR 2000-514584 [46]; 2001-514422 [56]

AB WO 200041548 A UPAB: 20021120

NOVELTY - A new method for lowering plasma **glucagon** comprises administering a compound (C1) selected from **exendin**, an **exendin** agonist, a modified **exendin** or a modified **exendin** agonist.

ACTIVITY - Antidiabetic; dermatological.

MECHANISM OF ACTION - The compounds lower plasma **glucagon** level.

The safety, tolerability, and efficacy of synthetic **exendin** -4 was evaluated in 8 male non-insulin using patients with type 2 diabetes who had discontinued other antidiabetic therapy for a minimum of 7 days. Each patient received subcutaneous (SC) injections of placebo (PBO) and 0.1, 0.2, and 0.3 micro g/kg **exendin**-4 48 hours apart in a single-blind, dose-rising, placebo controlled crossover design. Five patients also received a 0.4 micro g/kg dose. Plasma glucose, insulin and **glucagon** concentrations were assessed during fasting and in response to a 7 Kcal/kg Sustacal (RTM) challenge administered at the time of **exendin**-4/PBO injection. Gastric emptying was evaluated by measuring serum acetaminophen concentrations following a 20 mg/kg oral dose of liquid acetaminophen administered with the Sustacal (RTM).

No safety issues were identified based upon reported adverse events, EKG (undefined) and safety lab monitoring. Doses of 0.3 and 0.4 micro g/kg elicited a dose-dependent increase in nausea. Vomiting occurred at the highest dose.

Plasma glucose concentrations were reduced in all doses of **exendin**-4 compared to PBO although insulin concentrations were not significantly different. The 8 hour mean plus or minus SE changes in plasma glucose AUC (undefined) from baseline were +391 plus or minus 187, -263 plus or minus 108, -247 plus or minus 64, -336 plus or minus 139, and -328 plus or minus 70 (mg)(hr)/dL for the PBO, 0.1, 0.2, 0.3, and 0.4 micro g/kg doses respectively. The 3 hour changes in plasma **glucagon** were +128.0 plus or minus 19.2, -5.6 plus or minus 10.5, -29.4 plus or minus 18.6, -40.5 plus or minus 24.5, and +6.9 plus or minus 38.6 (pg)(hr)/mL respectively. The gastric emptying rate was slowed in all doses and the mean total absorbed acetaminophen over 6 hours was reduced by 51%, 50%, 57% and 79% compared to PBO for 0.1, 0.2, 0.3, and 0.4 micro g/kg doses respectively.

In summary, SC injection of **exendin**-4 to patients identified no safety issues, was tolerated at doses at most 0.3 micro g/kg, reduced plasma glucose and **glucagon** and slowed the rate of gastric emptying.

USE - The method is useful for lowering plasma **glucagon** in subjects, preferably humans, suffering from necrolytic erythema or **glucagonoma** (claimed). The method is also useful for treating hyperglucagonemia and other conditions that would benefit from reduced **glucagon** levels and/or suppression of **glucagon**, e.g. type 1 and type 2 diabetes.

Dwg.0/6

=> dup rem 1432

DUPLICATE IS NOT AVAILABLE IN 'DGENE, ADISINSIGHT, DRUGUPDATES, PHARMAML, ADISNEWS, PHAR, BIOCOMMERCE, FEDRIP, GENBANK'.

ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE

PROCESSING IS APPROXIMATELY 87% COMPLETE FOR L432

PROCESSING COMPLETED FOR L432

L474 737 DUP REM L432 (506 DUPLICATES REMOVED)

=> s 1474 (s) (glucagon (w) secretion)

L475 396 S L474

PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH

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L481 27 S L474  
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TOTAL FOR ALL FILES  
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 The previous command name entered was not recognized by the system.  
 For a list of commands available to you in the current file, enter  
 "HELP COMMANDS" at an arrow prompt (=>).

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 L557 2 FILE DGENE  
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TOTAL FOR ALL FILES

L636 46 L474 AND (GLUCAGON (W) SECRETION)

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NO VALID FORMATS ENTERED FOR FILE 'ADISINSIGHT'

In a multifile environment, each file must have at least one valid format requested. Refer to file specific help messages or the STNGUIDE file for information on formats available in individual files.

REENTER DISPLAY FORMAT FOR ALL FILES (FILEDEFAULT):so abs ti au

L636 ANSWER 1 OF 46 DGENE (C) 2003 THOMSON DERWENT

AN AAB85927 peptide DGENE  
AB The invention is directed towards the amelioration of organ tissue injury caused by reperfusion of blood flow after ischemia. The method involves administering a composition containing a compound which binds to a receptor for **glucagon**-like peptide-1 (GLP-1) in a carrier. GLP-1 effectively enhances peripheral glucose uptake without inducing dangerous hypoglycemia. GLP-1 strongly suppresses **glucagon secretion**, independent of its insuliniotropic action and powerfully reduces plasma free fatty acid (FFA) level having major toxic mechanism during myocardial ischemia, substantially more than can be accomplished with insulin. The method is without side effects normally attendant with therapies presently available. GLP-1 suppresses paracrine by intra-islet release of insulin or somatostatin. GLP-1 is unique in its capacity to simultaneously stimulate insulin secretion and inhibit **glucagon** release. The present sequence represents a gila monster venom **exendin 4** peptide fragment, homologous to a mammalian GLP-1 peptide fragment.  
TI Use of **glucagon**-like peptide-one for amelioration of organ tissue e.g. myocardium, injury after ischemia -  
IN Coolidge T R; Ehlers M R W

L636 ANSWER 2 OF 46 DGENE (C) 2003 THOMSON DERWENT

AN AAB85926 peptide DGENE  
AB The invention is directed towards the amelioration of organ tissue injury caused by reperfusion of blood flow after ischemia. The method involves administering a composition containing a compound which binds to a receptor for **glucagon**-like peptide-1 (GLP-1) in a carrier. GLP-1 effectively enhances peripheral glucose uptake without inducing dangerous hypoglycemia. GLP-1 strongly suppresses **glucagon secretion**, independent of its insuliniotropic action and powerfully reduces plasma free fatty acid (FFA) level having major toxic mechanism during myocardial ischemia, substantially more than can be accomplished with insulin. The method is without side effects normally attendant with therapies presently available. GLP-1 suppresses paracrine by intra-islet release of insulin or somatostatin. GLP-1 is unique in its capacity to simultaneously stimulate insulin secretion and inhibit **glucagon** release. The present sequence represents a gila monster venom **exendin 4** peptide fragment, homologous to a mammalian GLP-1 peptide fragment.  
TI Use of **glucagon**-like peptide-one for amelioration of organ tissue e.g. myocardium, injury after ischemia -  
IN Coolidge T R; Ehlers M R W

L636 ANSWER 3 OF 46 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

SO Biomedical Research (Tokyo), (December, 2002) Vol. 22, No. 6, pp. 295-297.  
print.  
ISSN: 0388-6107.

AB **Glucagon**-like peptide-1 (GLP-1) is known to lower blood glucose level, the effect depending on the stimulation of insulin and the inhibition of **glucagon secretion**. Due to the rapid inactivation of GLP-1 by dipeptidyl peptidase-IV (DPP-IV), its biological action is very short. Hence, we investigated the effect on rat entero-insular axis of **exendin-4**, a DPP-IV-resistant agonist of GPL-1 receptors. As expected the bolus administration of **exendin-4** (12 nmol/kg) increased the plasma concentration of insulin and decreased the blood levels of both **glucagon** and leptin in normal rats; however, GLP-1 raised glycemia. **Exendin-4** did not evoke any effect in rats bearing enucleated-regenerated adrenals deprived of medullary tissue, thereby suggesting that its stimulating effect of entero-insular axis occurs via an indirect mechanism probably involving medullary catecholamines. Catecholamines are potent stimulator of hepatic glycogenolysis, and this may tentatively explain the hyperglycemizing effect of **exendin-4**.

TI **Exendin-4**, a GLP-1 receptor agonist, stimulates



entero-insular axis in the rat, through a mechanism involving adrenal medulla.

AU Malendowicz, Ludwik K.; Nowak, Krzysztof W.; Zyterska, Agnieszka; Nussdorfer, Gastone G. (1); Macchi, Carlo; Nowak, Magdalena

L636 ANSWER 4 OF 46 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
SO Diabetes, (June, 2001) Vol. 50, No. Supplement 2, pp. A313. print.  
Meeting Info.: 61st Scientific Sessions of the American Diabetes Association Philadelphia, Pennsylvania, USA June 22-26, 2001.  
ISSN: 0012-1797.

TI Effect of **exendin-4** on **glucagon secretion** in lean and obese Zucker (ZDF) rats.

AU Parkes, David (1); Gedulin, Bronislava (1); Smith, Pamela (1); Young, Andrew (1)

L636 ANSWER 5 OF 46 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
SO Experimental and Clinical Endocrinology & Diabetes, (2001) Vol. 109, No. Suppl. 2, pp. S288-S303. print.  
ISSN: 0947-7349.

AB The search for intestinal factors regulating the endocrine secretion of the pancreas started soon after the discovery of secretin, i.e. nearly 100 years ago. Insulinotropic factors of the gut released by nutrients and stimulating insulin secretion in physiological concentrations in the presence of elevated blood glucose levels have been named incretins. Of the known gut hormones only gastric inhibitory polypeptide (GIP) and **glucagon-like polypeptide-1 (GLP-1 (7-36) amide)** fulfill this definition. - The incretin effect (i.e. the ratio between the integrated insulin response to an oral glucose load and an isoglycaemic intravenous glucose infusion) is markedly diminished in patients with type 2 diabetes mellitus, while the plasma levels of GIP and GLP-1 and their responses to nutrients are in the normal range. Therefore, a reduced responsiveness of the islet B-cells to incretins has been postulated. This insensitivity of the diabetic B-cells towards incretins can be overcome by supraphysiological (pharmacological) concentrations of GLP-1 (7-36), however not of GIP. Accordingly, fasting and postprandial glucose levels can be normalized in patients with type 2 diabetes by infusions of GLP-1 (7-36). Further studies revealed that this is partially due to the fact that GLP-1 (7-36) - in addition to its insulinotropic effect - also inhibits **glucagon secretion** and delays gastric emptying. These three antidiabetic effects qualify GLP-1 (7-36) as an interesting therapeutic tool, mainly for type 2 diabetes. However, because of its short plasma half life time natural GLP-1 (7-36) is not suitable for subcutaneous application. At present methods are being developed to improve the pharmacokinetics of GLP-1 by inhibition of the cleaving enzyme dipeptidyl peptidase IV (DPP-IV) or by synthesis of DPP-IV resistant GLP-1 analogues. Also naturally occurring GLP-1 analogues (for instance **exendin-4**) with a much longer half life time than GLP-1 (7-36) are being tested. - Thus, after 100 years of speculations and experimentations, incretins and their analogues are emerging as new antidiabetic drugs.

TI The entero-insular axis in type 2 diabetes - incretins as therapeutic agents.

AU Creutzfeldt, W. (1)

L636 ANSWER 6 OF 46 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
SO European Journal of Endocrinology, (December, 2000) Vol. 143, No. 6, pp. 717-725. print.  
ISSN: 0804-4643.

AB **Glucagon-like peptide-1 (GLP-1)** is a gut hormone synthesized by post-translational processing in intestinal L-cells, and it is released in response to food ingestion. GLP-1 stimulates insulin secretion during hyperglycemia, suppresses **glucagon secretion**, stimulates (pro)-insulin biosynthesis and decreases the rate of gastric emptying and acid secretion. GLP-1 has also been shown to have a pro-satiety effect. In addition, it has been demonstrated that a long-term

infusion with GLP-1, or **exendin-4**, a long-acting analog of human GLP-1, increases beta-cell mass in rats. In conclusion, GLP-1 appears to regulate plasma glucose levels via various and independent mechanisms. GLP-1 is an excellent candidate option for the treatment of patients with type 2 diabetes mellitus.

TI **Glucagon-like peptide-1: A major regulator of pancreatic beta-cell function.**

AU Perfetti, Riccardo (1); Merkel, Patricia

L636 ANSWER 7 OF 46 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

SO Diabetes, (1999) Vol. 48, No. SUPPL. 1, pp. A198.

Meeting Info.: 59th Scientific Sessions of the American Diabetes Association San Diego, California, USA June 19-22, 1999 American Diabetes Association

. ISSN: 0012-1797.

TI **Exendin-4 (AC2993) decreases glucagon secretion during hyperglycemic clamps in Diabetic Fatty Zucker rats.**

AU Gedulin, Bronislava (1); Jodka, Lynne (1); Hoyt, Julie (1)

L636 ANSWER 8 OF 46 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

SO Diabetes, (Feb., 1998) Vol. 47, No. 2, pp. 159-169.

ISSN: 0012-1797.

AB Proglucagon contains the sequence of two **glucagon-like peptides**, GLP-1 and GLP-2, secreted from enteroendocrine cells of the small and large intestine. GLP-1 lowers blood glucose in both NIDDM and IDDM patients and may be therapeutically useful for treatment of patients with diabetes. GLP-1 regulates blood glucose via stimulation of glucose-dependent insulin secretion, inhibition of gastric emptying, and inhibition of **glucagon secretion**. GLP-1 may also regulate glycogen synthesis in adipose tissue and muscle; however, the mechanism for these peripheral effects remains unclear. GLP-1 is produced in the brain, and intracerebroventricular GLP-1 in rodents is a potent inhibitor of food and water intake. The short duration of action of GLP-1 may be accounted for in part by the enzyme dipeptidyl peptidase 4 (DPP-IV), which cleaves GLP-1 at the NH<sub>2</sub>-terminus; hence GLP-1 analogs or the lizard peptide **exendin-4** that are resistant to DPP-IV cleavage may be more potent GLP-1 molecules in vivo. GLP-2 has recently been shown to display intestinal growth factor activity in rodents, raising the possibility that GLP-2 may be therapeutically useful for enhancement of mucosal regeneration in patients with intestinal disease. This review discusses recent advances in our understanding of the biological activity of the **glucagon-like peptides**.

TI **Glucagon-like peptides.**

AU Drucker, Daniel J. (1)

L636 ANSWER 9 OF 46 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

SO Journal of Biological Chemistry, (1997) Vol. 272, No. 7, pp. 4108-4115.

ISSN: 0021-9258.

AB **Glucagon-like peptide 1 stimulates insulin secretion and inhibits glucagon secretion**, gastric emptying, and feeding, suggesting it may be biologically useful for the treatment of diabetes. A lizard **glucagon-like peptide 1 (GLP-1)-related peptide, exendin 4**, binds to the GLP-1 receptor and mimics the actions of GLP-1 in vivo. To determine the genetic relationship between **exendin 4** and GLP-1, we analyzed the structure and expression of pancreatic and intestinal proglucagon mRNAs in the reptile *Heloderma suspectum*. Two different proglucagon cDNAs (lizard proglucagon I (LPI) and lizard proglucagon II (LPII)), with unique 3'-untranslated regions were identified. Two LPI mRNA transcripts, approx 1.6 and 2.1 kilobases, encoded **glucagon** and GLP-1 but not GLP-2 and were restricted in expression to the pancreas. In contrast, a 1.1-kilobase LPII mRNA transcript, encoding **glucagon**, GLP-1, and GLP-2 utilized a different 3'-untranslated region and was expressed in both pancreas and intestine. Lizard proglucagon mRNA transcripts were not

detectable by reverse transcription-polymerase chain reaction or Northern blotting in salivary gland. A single class of lizard salivary gland proexendin cDNAs encoded the sequence of **exendin 4** and a 45-amino acid **exendin** NH-2-terminal peptide. **Exendin** mRNA transcripts were expressed in the salivary gland, but not pancreas or intestine. These data demonstrate that GLP-1 and **exendin 4** represent related yet distinct peptides encoded by different genes in the lizard.

TI Tissue-specific expression of unique mRNAs that encode proglucagon-derived peptides or **exendin 4** in the lizard.

AU Chen, Yuqing E.; Drucker, Daniel J. (1)

L636 ANSWER 10 OF 46 CAPLUS COPYRIGHT 2003 ACS

SO Biomedical Research (2001), 22(6), 295-297  
CODEN: BRES5; ISSN: 0388-6107

AB **Glucagon**-like peptide-1 (GLP-1) is known to lower blood glucose level, the effect depending on the stimulation of insulin and the inhibition of **glucagon secretion**. Due to the rapid inactivation of GLP-1 by dipeptidyl peptidase-IV (DPP-IV), its biol. action is very short. Hence, we investigated the effect on rat entero-insular axis of **Exendin-4**, a DPP-IV-resistant agonist of GPL-1 receptors. As expected the bolus administration of **Exendin-4** (12 nmol/kg) increased the plasma concn. of insulin and decreased the blood levels of both **glucagon** and leptin in normal rats; however, GLP-1 raised glycemia. **Exendin-4** did not evoke any effect in rats bearing enucleated-regenerated adrenals deprived of medullary tissue, thereby suggesting that its stimulating effect of entero-insular axis occurs via an indirect mechanism probably involving medullary catecholamines. Catecholamines are potent stimulator of hepatic glycogenolysis, and this may tentatively explain the hyperglycemizing effect of **Exendin-4**.

TI **Exendin-4**, a GLP-1 receptor agonist, stimulates entero-insular axis in the rat, through a mechanism involving adrenal medulla

AU Malendowicz, Ludwik K.; Nowak, Krzysztof W.; Zyterska, Agnieszka; Nussdorfer, Gastone G.; Macchi, Carlo; Nowak, Magdalena

L636 ANSWER 11 OF 46 CAPLUS COPYRIGHT 2003 ACS

SO PCT Int. Appl., 96 pp.  
CODEN: PIXXD2

AB Methods are provided for use of an **exendin**, an **exendin** agonist, or a modified **exendin** or **exendin** agonist having an **exendin** or **exendin** agonist linked to one or more polyethylene glycol polymers, for example, for lowering **glucagon** levels and/or suppressing **glucagon secretion** in a subject. These methods are useful in treating hyperglucagonemia and other conditions that would be benefited by lowering plasma **glucagon** or suppressing **glucagon secretion**.

TI Methods using an **exendin** or related substance for **glucagon** suppression

IN Young, Andrew; Gedulin, Bronislava

L636 ANSWER 12 OF 46 SCISEARCH COPYRIGHT 2003 ISI (R)

SO DIABETES, (MAY 2000) Vol. 49, Supp. [1], pp. 460-460.  
Publisher: AMER DIABETES ASSOC, 1660 DUKE ST, ALEXANDRIA, VA 22314.  
ISSN: 0012-1797.

TI Dose-response for inhibition of **glucagon secretion** and gastric emptying by synthetic **exendin-4** (AC2993) in subjects with type 2 diabetes

AU Kolterman O (Reprint); Gottlieb A; Prickett K; Gaines E; Young A

L636 ANSWER 13 OF 46 SCISEARCH COPYRIGHT 2003 ISI (R)

SO DIABETES, (JUN 1999) Vol. 48, Supp. [1], pp. 864-864.  
Publisher: AMER DIABETES ASSOC, 1660 DUKE ST, ALEXANDRIA, VA 22314.

ISSN: 0012-1797.

- TI **Exendin-4** (AC2993) decreases **glucagon secretion** during hyperglycemic clamps in Diabetic Fatty Zucker rats
- AU Gedulin B (Reprint); Jodka L; Hoyt J
- L636 ANSWER 14 OF 46 Elsevier BIOBASE COPYRIGHT 2003 Elsevier Science B.V.  
SO Experimental and Clinical Endocrinology and Diabetes, (2001), 109/SUPPL. 2 (S288-S303), 122 reference(s)  
CODEN: ECEDFQ ISSN: 0947-7349
- AB The search for intestinal factors regulating the endocrine secretion of the pancreas started soon after the discovery of secretin, i.e. nearly 100 years ago. Insulinotropic factors of the gut released by nutrients and stimulating insulin secretion in physiological concentrations in the presence of elevated blood glucose levels have been named incretins. Of the known gut hormones only gastric inhibitory polypeptide (GIP) and **glucagon-like polypeptide-1** (GLP-1 [7 - 36] amide) fulfill this definition. - The incretin effect (i.e. the ratio between the integrated insulin response to an oral glucose load and an isoglycaemic intravenous glucose infusion) is markedly diminished in patients with type 2 diabetes mellitus, while the plasma levels of GIP and GLP-1 and their responses to nutrients are in the normal range. Therefore, a reduced responsiveness of the islet B-cells to incretins has been postulated. This insensitivity of the diabetic B-cells towards incretins can be overcome by supraphysiological (pharmacological) concentrations of GLP-1 [7 - 36], however not of GIP. Accordingly, fasting and postprandial glucose levels can be normalized in patients with type 2 diabetes by infusions of GLP-1 [7 - 36]. Further studies revealed that this is partially due to the fact that GLP-1 [7 - 36] - in addition to its insulinotropic effect - also inhibits **glucagon secretion** and delays gastric emptying. These three antidiabetic effects qualify GLP-1 [7 - 36] as an interesting therapeutic tool, mainly for type 2 diabetes. However, because of its short plasma half life time natural GLP-1 [7 - 36] is not suitable for subcutaneous application. At present methods are being developed to improve the pharmacokinetics of GLP-1 by inhibition of the cleaving enzyme dipeptidyl peptidase IV (DPP-IV) or by synthesis of DPP-IV resistant GLP-1 analogues. Also naturally occurring GLP-1 analogues (for instance **exendin-4**) with a much longer half life time than GLP-1 [7 - 36] are being tested. - Thus, after 100 years of speculations and experimentations, incretins and their analogues are emerging as new antidiabetic drugs.
- TI The entero-insular axis in type 2 diabetes - Incretins as therapeutic agents
- AU Creutzfeldt W.
- L636 ANSWER 15 OF 46 DRUGU COPYRIGHT 2003 THOMSON DERWENT  
SO Diabetologia (45, Suppl. 2, A214, 2002)  
CODEN: DBTGAI ISSN: 0012-186X
- AV Department of Physiology, Clinica Puerta de Hierro, Universidad Autonoma de Madrid, Madrid, Spain.
- AN 2002-37701 DRUGU P E
- AB In-vitro, **exendin-4** (Amylin) directly inhibited alpha-cell secretion in perfused rat pancreas. **Exendin-4** is an insulinotropic peptide structurally analogous to GLP-1. (conference abstract: 38th Annual Meeting of the European Association for the Study of Diabetes, Budapest, Hungary, 2002).
- ABEX **Exendin-4** (1 nM) appreciably potentiated insulin and somatostatin responses to arginine (10 mM) and appreciably reduced **glucagon** response to this aminoacid by about 3x. **Exendin-4** (1 nM) appreciably reduced the increase in **glucagon secretion** induced by abrupt glucose depletion (from 11 to 3.2 mM) but had no effect on the dramatic decrease in insulin release or the slight decrease in somatostatin release under these conditions. (E42/JM)
- TI Evidence for a direct inhibitory effect of **exendin-4** on alpha cell secretion.

- AU Rodriguez Gallardo J; Egido E M; Gutierrez E; Garcia P; Silvestre R A; Marco J
- L636 ANSWER 16 OF 46 DRUGU COPYRIGHT 2003 THOMSON DERWENT  
 SO Diabetes (51, Suppl. 2, A339-A340, 2002)  
 CODEN: DIAEAZ ISSN: 0012-1797  
 AV No Reprint Address.  
 AN 2002-31847 DRUGU P E  
 AB The effects of **exendin-4** and **glucagon**-like peptide (GLP)-1 were compared in isolated perfused pig pancreas preparations. Both peptides slightly reduced vascular resistance but neither affected pancreas exocrine excretion. Both peptides stimulated insulin (up to 203% and 309% respectively) and somatostatin (208% and 105%) at 0.01-10 nM and both inhibited **glucagon secretion**. It is concluded that **exendin-4** and GLP-1 have identical endocrine effects and similar potency in the pig pancreas. (conference abstract: 62nd Scientific Sessions of the American Diabetes Association, San Francisco, California, USA, 2002). (No EX).  
 ABEX (E33/JB)  
 TI **Exendin-4** and **glucagon**-like peptide-1 have similar effects in isolated perfused porcine pancreas.  
 AU Hansen L; Nielsen J Z; Holst J J
- L636 ANSWER 17 OF 46 DRUGU COPYRIGHT 2003 THOMSON DERWENT  
 SO Diabetes (50, Suppl. 2, A313, 2001)  
 CODEN: DIAEAZ ISSN: 0012-1797  
 AV No reprint address.  
 AN 2002-03605 DRUGU P E  
 AB The effect of i.v. **exendin-4** on **glucagon** synthesis was investigated in a rat model of type 2 diabetes. **Exendin-4** suppressed **glucagon secretion** in lean and obese diabetic Zucker rats via a mechanism distinct from elevation of plasma glucose. The results support the use of **exendin-4** in diabetic conditions where excessive **glucagon secretion** contributes to fasting hyperglycemia. (conference abstract: 61st Scientific Sessions of the American Diabetes Association, Philadelphia, Pennsylvania, USA, 2001).  
 ABEX Methods Lean and obese (diabetic) Zucker rats were administered i.v. **exendin-4** (2.6 ug/hr) or saline for 30 min. Results In obese rats treated with **exendin-4**, **glucagon** decreased from 94 to 54 pM despite a decrease in plasma glucose. In lean rats, **exendin-4** decreased **glucagon** from 61 to 44 pM at 30 min accompanied by a decrease in plasma glucose. (E97)  
 TI Effect of **exendin-4** on **glucagon** synthesis in lean and obese Zucker (ZDF) rats.  
 AU Parkes D; Gedulin B; Smith P; Young A
- L636 ANSWER 18 OF 46 DRUGU COPYRIGHT 2003 THOMSON DERWENT  
 SO Exp.Clin.Endocrinol.Diabetes (109, Suppl. 2, S288-S203, 2001) 2 Fig. 1 Tab. 122 Ref.  
 CODEN: ECEDF ISSN: 0947-7349  
 AV Zentrum Innere Medizin, Klinikum der Universitaet, Robert- Koch-Str. 40 D-37075 Goettingen, Germany.  
 AN 2001-26811 DRUGU T E  
 AB Incretins as therapeutics agents for type 2 diabetes is reviewed with specific reference to **glucagon**-like polypeptide-1 (GLP-1 [7-36] amide).  
 ABEX Insulinotropic factors of the gut release by nutrients and stimulating insulin secretion in physiological concentrations in the presence of elevated blood glucose levels had been named incretins. Of the known gut hormones only gastric inhibitory polypeptide (GIP) and **glucagon**-like polypeptide-1 (GLP-1 [7-36] amide) fulfill this definition. The incretin effect (i.e. the ratio between the integrated insulin response to an oral glucose load and an isoglycemic i.v. glucose infusion) is

markedly diminished in patients with type 2 diabetes mellitus, while the plasma levels of GIP and GLP-1 and their response to nutrients are in the normal range. Therefore a reduced responsiveness to the islet B cells to incretins has been postulated, This insensitivity of the diabetic B-cells towards incretins can be overcome by supraphysiological concentrations of GLP-1 [7-36], however not of GIP. Accordingly, fasting and postprandial glucose levels can be normalized in patients with type 2 diabetes by infusions of GLP-1 [7-36]. GLP-1 [7-36] also inhibits **glucagon secretion** and delays gastric emptying. Because of its short plasma half-life time natural GLP-1 [7-36] is not suitable for s.c. application. Naturally occurring GLP-1 analogs (e.g. **exendin-4**) with a much longer half-life time than GLP-1 [7-36] are being tested. (E98)

TI The entero-insular axis in type 2 diabetes - incretins as therapeutic agents.

AU Creutzfeldt W

L636 ANSWER 19 OF 46 DRUGU COPYRIGHT 2003 THOMSON DERWENT

SO Diabetes (49, Suppl. 1, A114, 2000)  
CODEN: DIAEAZ ISSN: 0012-1797

AV No Reprint Address.

AN 2000-40616 DRUGU T E

AB S.c. synthetic **exendin-4** (AC-2993) inhibited postprandial plasma glucose and **glucagon** elevations and gastric emptying in 14 patients with type-2 diabetes. Each of these effects demonstrated similar dose potency. (conference abstract: 60th Scientific Sessions of the American Diabetes Association, San Antonio, Texas, USA, 2000).

ABEX Methods Patients (mean age 55 yr) received single injections of AC-2993 (0.01, 0.02, 0.05 or 0.1 ug/kg) or placebo on separate days following an overnight fast. Injections were given immediately prior to ingestion of a standard Sustacal meal (7 kcal/kg). Gastric emptying was evaluated by giving liquid phenazone (20 mg/kg) along with the meal and monitoring phenazone appearance in plasma. Results AC-2993 inhibited postprandial glucose and **glucagon** elevations and gastric-emptying in a dose-dependent manner with ED50 values of 0.038, 0.017 and 0.048 ug/kg, respectively. (E42/JM)

TI Dose-response for inhibition of **glucagon secretion** and gastric emptying by synthetic **exendin-4** (AC2993) in subjects with type 2 diabetes.

AU Kolterman O; Gottlieb A; Prickett K; Gaines E; Young A

L636 ANSWER 20 OF 46 DRUGU COPYRIGHT 2003 THOMSON DERWENT

SO Diabetes (48, Suppl. 1, A199, 1999)  
CODEN: DIAEAZ ISSN: 0012-1797

AV No Reprint Address.

AN 1999-28905 DRUGU P E

AB Effects of i.v. infusion of **exendin-4** (AC-2993), a 39 amino acid peptide isolated from salivary secretions of the Gila monster (*Heloderma suspectum*), with antidiabetic actions similar to those of GLP-1, on **glucagon secretion** were investigated in anesthetized male diabetic fatty Zucker (ZDF) rats. **Exendin-4** exhibited a glucagonostatic effect in ZDF rats during hyperglycemic clamp studies, an effect which if present in diabetic humans, could be of potential therapeutic benefit. (conference abstract: 59th Annual Scientific Sessions of the American Diabetes Association, San Diego, California, USA, 1999).

ABEX Using an hyperinsulinemic hyperglycemic clamp protocol, those factors tending to influence **glucagon secretion** were held constant. Plasma glucose was clamped at about 34 mM 60 min before beginning infusions of saline or **exendin-4** (0.21 +/- 2.1 ug/ml/hr). Plasma **glucagon** concentration measured prior to these infusions were similar in both groups (306 vs. 252 pM, respectively). Mean plasma **glucagon** concentration in **exendin-4** infused rats was nearly half of that in

saline-infused rats in the final 60 min of the clamp (165 vs. 2984 pM, respectively). The hyperglycemic clamp protocol also enabled the measurement of insulin sensitivity. Glucose infusion rate during the clamp was increased by 111% in **exendin-4**-treated vs. control rats. (E54/RSV)

TI **Exendin-4** (AC2993) decreases **glucagon-secretion** during hyperglycemic clamps in diabetic fatty Zucker rats.

AU Gedulin B; Jodka L; Hoyt J

L636 ANSWER 21 OF 46 USPATFULL

AB Methods for treating gestational diabetes which comprise administration of an effective amount of an **exendin** or an **exendin** agonist, alone or in conjunction with other compounds or compositions that lower blood glucose levels.

TI Use of **exendins** and agonists thereof for the treatment of gestational diabetes mellitus

IN Hiles, Richard A., San Diego, CA, United States  
Prickett, Kathryn S., San Diego, CA, United States

L636 ANSWER 22 OF 46 USPATFULL

AB Dipeptidyl peptidase IV (DP 4) inhibiting compounds are provided having the formula ##STR1##

where n is 0 or 1; X is H or CN;

Y is N, NH or O;

Z is CH.sub.2 when Y is O or N--H, with Y--Z forming a single bond, and Z is CH when Y is N, with Y--Z forming a double bond;

and wherein R.sup.1, R.sup.2, R.sup.3 and R.sup.4 are as described herein.

A method is also provided for treating diabetes and related diseases, especially Type II diabetes, and other diseases as set out herein, employing such DP 4 inhibitor or a combination of such DP 4 inhibitor and one or more of another antidiabetic agent such as metformin, glyburide, troglitazone, pioglitazone, rosiglitazone and/or insulin and/or one or more of a hypolipidemic agent and/or anti-obesity agent and/or other therapeutic agent.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI 2,1-Oxazoline and 1,2-pyrazoline-based inhibitors of dipeptidyl peptidase IV and method

IN Sulsky, Richard B., West Trenton, NJ, UNITED STATES  
Robl, Jeffrey A., Newtown, PA, UNITED STATES

L636 ANSWER 23 OF 46 USPATFULL

AB Individuals in need of treatment of ischemia-related reperfusion are treated, preferably intravenously, with a composition which includes a compound which binds to a receptor for the **glucagon**-like peptide-1. The invention relates to both the method and compositions for such treatment.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Metabolic intervention with GLP-1 to improve the function of ischemic and reperfused skeletal muscle tissue

IN Coolidge, Thomas R., Falls Village, CT, UNITED STATES  
Ehlers, Mario R.W., Lincoln, NE, UNITED STATES

L636 ANSWER 24 OF 46 USPATFULL

AB Methods for treating conditions or disorders which can be alleviated by reducing food intake are disclosed which comprise administration of an

effective amount of an exendin or an exendin agonist, alone or in conjunction with other compounds or compositions that affect satiety. The methods are useful for treating conditions or disorders, including obesity, Type II diabetes, eating disorders, and insulin-resistance syndrome. The methods are also useful for lowering the plasma glucose level, lowering the plasma lipid level, reducing the cardiac risk, reducing the appetite, and reducing the weight of subjects. Pharmaceutical compositions for use in the methods of the invention are also disclosed.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI USE OF EXENDINS AND AGONISTS THEREOF FOR THE REDUCTION OF FOOD INTAKE  
IN BEELEY, NIGEL ROBERT ARNOLD, SOLANA BEACH, CA, UNITED STATES  
PRICKETT, KATHRYN S., SAN DIEGO, CA, UNITED STATES  
BHAVSAR, SUNIL, SAN DIEGO, CA, UNITED STATES

L636 ANSWER 25 OF 46 USPATFULL

AB The effects of GLP-2 are enhanced using a GLP-1 activity inhibitor. For medical use to treat or inhibit the onset of medical conditions, disorder or diseases for which treatment with GLP-2 is indicated, the present invention provides a pharmaceutical combination comprising a GLP-2 activity enhancer, and a GLP-1 activity inhibitor. The combination is useful particularly to treat gastrointestinal conditions such as small bowel syndrome, mucositis and Crohn's disease, and to suppress appetite, for instance to treat obesity.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Enhancement of GLP-2 activity  
IN Drucker, Daniel J., Toronto, CANADA  
Lovshin, Julie Ann Louise, Toronto, CANADA

L636 ANSWER 26 OF 46 USPATFULL

AB Individuals in need of treatment of ischemia-related reperfusion are treated, preferably intravenously, with a composition which includes a compound which binds to a receptor for the **glucagon**-like peptide-1. The invention relates to both the method and compositions for such treatment.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Metabolic intervention with GLP-1 to improve the function of ischemic and reperfused tissue  
IN Coolidge, Thomas R., Falls Village, CT, UNITED STATES  
Ehlers, Mario R.W., Lincoln, NE, UNITED STATES

L636 ANSWER 27 OF 46 USPATFULL

AB Modified insulintropic peptides are disclosed. The modified insulintropic peptides are capable of forming a peptidase stabilized insulintropic peptide. The modified insulintropic peptides are capable of forming covalent bonds with one or more blood components to form a conjugate. The conjugates may be formed in vivo or ex vivo. The modified peptides are administered to treat humans with diabetes and other related diseases.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Long lasting insulinoptropic peptides  
IN Bridon, Dominique P., Outremont, CANADA  
L'Archeveque, Benoit, Leval, CANADA  
Ezrin, Alan M., Moraga, CA, UNITED STATES  
Holmes, Darren L., Montreal, CANADA  
Leblanc, Anouk, Montreal, CANADA  
St. Pierre, Serge, Ile Bizard, CANADA

L636 ANSWER 28 OF 46 USPATFULL

AB Compositions and methods using same for the treatment of diabetes its sequelae and pre-diabetic conditions are provided. Invention



compositions include the anti-diabetic agent metformin, and bioavailable sources of one or more of chromium, vanadium and magnesium. Also provided are pharmaceutical agents containing invention compositions and methods for administering such agents.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Metformin-containing compositions for the treatment of diabetes  
IN Fine, Stuart A., Northbrook, IL, United States  
Kinsella, Kevin J., La Jolla, CA, United States

L636 ANSWER 29 OF 46 USPTAFULL

AB Dipeptidyl peptidase IV (DP 4) inhibiting compounds are provided having the formula ##STR1##

where x is 0 or 1 and y is 0 or 1 (provided that x=1 when y=0 and x=0 when y=1);

n is 0 or 1; X is H or CN;

and wherein R<sup>sup.1</sup>, R<sup>sup.2</sup>, R<sup>sup.3</sup> and R<sup>sup.4</sup> are as described herein.

A method is also provided for treating diabetes and related diseases, especially Type II diabetes, and other diseases as set out herein, employing such DP 4 inhibitor or a combination of such DP 4 inhibitor and one or more of another antidiabetic agent such as metformin, glyburide, troglitazone, pioglitazone, rosiglitazone and/or insulin and/or one or more of a hypolipidemic agent and/or anti-obesity agent and/or other therapeutic agent.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Cyclopropyl-fused pyrrolidine-based inhibitors of dipeptidyl peptidase IV and method  
IN Robl, Jeffrey A., Newtown, PA, UNITED STATES  
Sulsky, Richard B., West Trenton, NJ, UNITED STATES  
Augeri, David J., Princeton, NJ, UNITED STATES  
Magnin, David R., Hamilton, NJ, UNITED STATES  
Hamann, Lawrence G., Cherry Hill, NJ, UNITED STATES  
Betebenner, David A., Lawrenceville, NJ, UNITED STATES

L636 ANSWER 30 OF 46 USPTAFULL

AB Since **glucagon**-like peptide-1 (GLP-1) is the most potent insulinotropic hormone known and has been shown to stimulate insulin secretion strongly in patients with type II diabetes, this invention uses GLP-1 or its biologically active analogues in .beta.-cell stimulatory tests in order to test .beta.-cell function in a simple way. The test provides information about insulin secretory capacity, is easy and reproducible and has insignificant side effects.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI GLP-1 as a diagnostic test to determine .beta.-cell function and the presence of the condition of IGT and type II diabetes  
IN Holst, J. J., Copenhagen, DENMARK  
Vilsboll, Tina, Hellerup, DENMARK

L636 ANSWER 31 OF 46 USPTAFULL

AB Modified insulinotropic peptides are disclosed. The modified insulinotropic peptides are capable of forming a peptidase stabilized insulinotropic peptide. The modified insulinotropic peptides are capable of forming covalent bonds with one or more blood components to form a conjugate. The conjugates may be formed in vivo or ex vivo. The modified peptides are administered to treat humans with diabetes and other related diseases.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Long lasting insulinotropic peptides  
IN Bridon, Dominique P., Outremont, Canada  
L'Archeveque, Benoit, Laval, Canada  
Ezrin, Alan M., Moraga, CA, United States  
Holmes, Darren L., Montreal, Canada  
Leblanc, Anouk, Montreal, Canada  
St. Pierre, Serge, Ile Bizard, Canada

L636 ANSWER 32 OF 46 USPATFULL

AB The present invention relates to a derivative of GLP-1 (7-C), wherein C is 35 or 36 which derivative has just one lipophilic substituent which is attached to the C-terminal amino acid residue.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Extendin derivatives  
IN Knudsen, Liselotte Bjerre, Valby, Denmark  
Huusfeldt, Per Olaf, Copenhagen K, Denmark  
Nielsen, Per Franklin, Vaerlose, Denmark  
Madsen, Kjeld, Vaerlose, Denmark

L636 ANSWER 33 OF 46 USPATFULL

AB Individuals in need of treatment of ischemia-related reperfusion are treated, preferably intravenously, with a composition which includes a compound which binds to a receptor for the **glucagon**-like peptide-1. The invention relates to both the method and compositions for such treatment.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Metabolic intervention with GLP-1 to improve the function of ischemic and reperfused tissue  
IN Coolidge, Thomas R., Falls Village, CT, United States  
Ehlers, Mario R. W., Lincoln, NE, United States

L636 ANSWER 34 OF 46 ADISCTI COPYRIGHT 2003 (ADIS)

SO Diabetologia (Oct 1, 2002), Vol. 45, pp. 1410-1415

TI Lack of effect of **exendin-4** and **glucagon**-like peptide-1-(7,36)-amide on insulin action in non-diabetic humans.  
ADIS TITLE: **Exendin 4**, **glucagon**-like peptide-1 (7,36) amide: pharmacodynamics.  
Effects on insulin action  
In volunteers.

AU Vella A; Shah P; Reed A S; Adkins A S; Basu R; et al.

L636 ANSWER 35 OF 46 ADISCTI COPYRIGHT 2003 (ADIS)

SO Diabetologia (Aug 1, 2002), Vol. 45 (Suppl. 2), pp. 44

TI 28 days of treatment with AC2993 (synthetic **exendin-4**) improved glycemic control in patients with type 2 diabetes concomitantly treated with metformin and sulfonylurea.

ADIS TITLE: **Exendin 4**: therapeutic use.

Type 2 diabetes mellitus

In obese patients receiving metformin and/or sulphonylureas.

AU Fineman M; Bicsak T; Shen L; Taylor K; Gaines E; et al.

L636 ANSWER 36 OF 46 ADISCTI COPYRIGHT 2003 (ADIS)

SO Diabetes (Jun 1, 2002), Vol. 51 (Suppl. 2), pp. 104 (plus poster)

TI Subcutaneous injection of AC2993 (synthetic **exendin-4**) lowered fasting glucose concentrations through suppression of **glucagon** and dose dependent insulinotropism in patients with type 2 diabetes.

ADIS TITLE: **Exendin 4**: pharmacodynamics.

Effects on fasting plasma glucose, serum insulin and plasma **glucagon** levels

In patients with type 2 diabetes mellitus receiving metformin and/or thiazolidinediones.

AU Kim D; Taylor K; Bicsak T; Wang Y; Aisporna M; et al.

- L636 ANSWER 37 OF 46 ADISCTI COPYRIGHT 2003 (ADIS)  
 SO Diabetes (Jun 1, 2002), Vol. 51 (Suppl. 2), pp. 84-85  
 TI **Exendin-4** reduces glycemic excursions after meals in insulin-treated diabetes.  
 ADIS TITLE: **Exendin 4**: pharmacodynamics.  
 Effects on postprandial gastric emptying  
 In patients with type 1 diabetes mellitus receiving insulin.  
 AU Dupre J; Behme M T; McDonald T J.
- L636 ANSWER 38 OF 46 ADISCTI COPYRIGHT 2003 (ADIS)  
 SO Diabetes (May 1, 2000), Vol. 49 (Suppl. 1), pp. 114 (plus poster)  
 TI Dose-response for inhibition of **glucagon secretion** and gastric emptying by synthetic **exendin-4** (AC2993) in subjects with type 2 diabetes.  
 ADIS TITLE: **Exendin-4**: pharmacodynamics.  
 Effects on **glucagon secretion** and gastric emptying  
 In patients with type 2 diabetes mellitus.  
 AU Kolterman O; Gottlieb A; Prickett K; Gaines E; Young A.
- L636 ANSWER 39 OF 46 ADISCTI COPYRIGHT 2003 (ADIS)  
 SO Diabetologia (Aug 1, 1999), Vol. 42 (Suppl. 1), pp. 41 (plus oral presentation)  
 TI AC2993 (synthetic **exendin-4**) lowered postprandial plasma glucose concentrations in people with type 2 diabetes.  
 ADIS TITLE: **Exendin-4**: pharmacodynamics.  
 Effect on postprandial glucose levels  
 In patients with type 2 diabetes mellitus.  
 AU Kolterman O; Fineman M; Gottlieb A; Petrella E; Prickett K; et al.
- L636 ANSWER 40 OF 46 ADISCTI COPYRIGHT 2003 (ADIS)  
 SO Diabetes (May 1, 1999), Vol. 48 (Suppl. 1), pp. 199  
 TI **Exendin-4** (AC993) decreases **glucagon secretion** during hyperglycemic clamps in diabetic fatty Zucker rats.  
 AU Gedulin B; Jodka L; Hoyt J.
- L636 ANSWER 41 OF 46 PROMT COPYRIGHT 2003 Gale Group  
 SO Marketletter, (30 Sep 2002) .  
 ISSN: 0951-3175.  
 AB In a ringing endorsement for Amylin Pharmaceuticals, Eli Lilly has licensed rights to the former's AC2993 diabetes drug in a deal which could be worth up to \$330 million in direct payments, including \$80 million in upfront nonrefundable fees, and stock purchases. Amylin's shares had put on nearly 12% to reach \$13.88 as this story went to press, while Lilly also posted a small gain to \$55.50.  
 THIS IS THE FULL TEXT: COPYRIGHT 2002 Marketletter Publications Ltd.  
 Subscription: 499.00 British pounds per year. Published weekly. 54-55 Wilton Road, London SW1V 1DE., United Kingdom  
 TI Amylin boosted as Lilly licenses diabetes drug in \$330M deal. (Brief Article)
- L636 ANSWER 42 OF 46 PROMT COPYRIGHT 2003 Gale Group  
 SO Marketletter, (24 Aug 1998) pp. N/A.  
 ISSN: 0951-3175.  
 AB Amylin of the USA has started clinical trials of its second drug candidate, **exendin-4**, in patients with type 2 diabetes. The drug is a synthetic version of a compound derived from the saliva of the Gila monster, a lizard native to the deserts of Arizona in the USA. The first Phase I study will be conducted in the UK and will look at escalating single doses of subcutaneous **exendin-4** in healthy volunteers. If the results are positive, proof-of-concept

studies in patients with type 2 diabetes could begin in 1999.

**Exendin-4** is similar in structure to **glucagon**

-like peptide-1, a hormone thought to be important in human glucose metabolism. Clinical data presented at an Eli Lilly meeting in Hamburg, Germany, last year, showed that GLP-1 produces a glucose-dependent stimulation of insulin secretion, an inhibition of **glucagon secretion**, increased rate of proinsulin synthesis and slowed gastric emptying. In addition, animal data has suggested that GLP-1 has insulin-like properties at target tissues (Marketletter May 5, 1997). Lilly is developing synthetic analogs of GLP-1, as the hormone itself has a plasma half-life which is too short to make a commercially-useful product. Amylin notes that **exendin-4** shares many of the properties of GLP-1 but offers a much longer biological duration of action. In animal tests, **exendin-4** stimulated secretion of insulin in hyperglycemia but not hypoglycemic conditions and also modulated gastric emptying. Most importantly, notes Amylin, **exendin-4** achieved a near-normalization of glucose control in an animal model of type 2 diabetes. Chronic administration to obese animals decreased food intake and led to a reduction in weight, suggesting **exendin-4** may also have a role to play in the management of obesity. Amylin's lead product, pramlintide, is currently in four pivotal trials in patients with both type 1 and type 2 diabetes (Marketletters passim).

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TI Amylin Starts Trials Of **Exendin-4** In Diabetes

L636 ANSWER 43 OF 46 COPYRIGHT 2003 Gale Group

SO Marketletter, (24 Aug 1998) .

ISSN: 0951-3175.

TI Amylin Starts Trials Of **Exendin-4** In Diabetes

L636 ANSWER 44 OF 46 ADISINSIGHT COPYRIGHT 2003 (ADIS)

SO Adis R&D Insight

L636 ANSWER 45 OF 46 PHARMAML COPYRIGHT 2003 MARKETLETTER

SO Marketletter September 30, 2002

AN 1665640 PHARMAML

TX In a ringing endorsement for Amylin Pharmaceuticals, Eli Lilly has licensed rights to the former's AC2993 diabetes drug in a deal which could be worth up to \$330 million in direct payments, including \$80 million in upfront nonrefundable fees, and stock purchases. Amylin's shares had put on nearly 12% to reach \$13.88 as this story went to press, while Lilly also posted a small gain to \$55.50.

AC2993 is a synthetic form of **exendin-4** which has a number of the biological activities of the hormone **glucagon** -like peptide-1, including the stimulation of insulin secretion in the presence of elevated blood glucose concentrations (but not during periods of low blood glucose concentrations), suppression of **glucagon secretion**, reduction of appetite and delay of food absorption.

However, Amylin's version, which is in Phase III trials, is much longer-acting than the native hormone. It is being developed as a fixed-dose injection in the first instance and could be filed for approval in this form as early as 2004. A key advantage over current therapies for type 2 diabetes is its low propensity to cause weight gain.

Lilly's interest in the compound stems from its own activities in diabetes, with a franchise that includes various forms of insulin and oral antidiabetic agents. Aside from the initial \$80 million license fee, the company will buy \$30 million-worth of Amylin stock at a premium to the current price, with a further commitment for additional payments of \$85 million on the attainment of certain developmental milestones and \$135 million on commercialization. Lilly may also make available a \$110

million convertible loan to fund a portion of Amylin's development and commercialization costs.

The two companies will share development and marketing costs for the product in the USA, with Lilly picking up 80% and 100% of these costs, respectively, outside this market. Profits will be shared in the USA, and Lilly will get 80% profit share in other countries.

Additionally, the companies have agreed that, for a limited period of time prior to the commercialization of AC2993, Amylin will co-promote Humatrope (somatropin), Lilly's recombinant human growth hormone product, in the USA.

TI Amylin boosted as Lilly licenses diabetes drug in \$330M deal

L636 ANSWER 46 OF 46 PHARMAML COPYRIGHT 2003 MARKETLETTER

SO Marketletter August 19, 1998

AN 1643252 PHARMAML

TX Amylin of the USA has started clinical trials of its second drug candidate, **exendin-4**, in patients with type 2 diabetes. The drug is a synthetic version of a compound derived from the saliva of the Gila monster, a lizard native to the deserts of Arizona in the USA.

The first Phase I study will be conducted in the UK and will look at escalating single doses of subcutaneous **exendin-4** in healthy volunteers. If the results are positive, proof-of-concept studies in patients with type 2 diabetes could begin in 1999.

**Exendin-4** is similar in structure to **glucagon**-like peptide-1, a hormone thought to be important in human glucose metabolism. Clinical data presented at an Eli Lilly meeting in Hamburg, Germany, last year, showed that GLP-1 produces a glucose-dependent stimulation of insulin secretion, an inhibition of **glucagon** secretion, increased rate of proinsulin synthesis and slowed gastric emptying. In addition, animal data has suggested that GLP-1 has insulin-like properties at target tissues (Marketletter May 5, 1997).

Lilly is developing synthetic analogs of GLP-1, as the hormone itself has a plasma half-life which is too short to make a commercially-useful product. Amylin notes that **exendin-4** shares many of the properties of GLP-1 but offers a much longer biological duration of action.

In animal tests, **exendin-4** stimulated secretion of insulin in hyperglycemia but not hypoglycemic conditions and also modulated gastric emptying. Most importantly, notes Amylin, **exendin-4** achieved a near-normalization of glucose control in an animal model of type 2 diabetes. Chronic administration to obese animals decreased food intake and led to a reduction in weight, suggesting **exendin-4** may also have a role to play in the management of obesity.

Amylin's lead product, pramlintide, is currently in four pivotal trials in patients with both type 1 and type 2 diabetes (Marketletters passim).

TI Amylin Starts Trials Of **Exendin-4** In Diabetes

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**Search Results - Record(s) 1 through 3 of 3 returned.**☐ 1. Document ID: US 20020010133 A1

L2: Entry 1 of 3

File: PGPB

Jan 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020010133

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020010133 A1

TITLE: METHOD FOR PREVENTING GASTRITIS USING AMYLIN OR AMYLIN AGONISTS

PUBLICATION-DATE: January 24, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
YOUNG, ANDREW A.	SAN DIEGO	CA	US	
GEDULIN, BRONISLAVA	SAN DIEGO	CA	US	
BEYNON, GARETH W.	BRIGHTWELL-CUM SOTWELL		UA	

US-CL-CURRENT: 514/12; 514/13, 514/14

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 2. Document ID: WO 9850059 A1

L2: Entry 2 of 3

File: EPAB

Nov 12, 1998

PUB-NO: WO009850059A1

DOCUMENT-IDENTIFIER: WO 9850059 A1

TITLE: METHOD FOR PREVENTING GASTRITIS USING AMYLIN OR AMYLIN AGONISTS

PUBN-DATE: November 12, 1998

## INVENTOR-INFORMATION:

NAME	COUNTRY
YOUNG, ANDREW	US
GEDULIN, BRONISLAVA	US
BEYNON, GARETH WYN	GB

INT-CL (IPC): A61 K 38/10; A61 K 38/16

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 3. Document ID: WO 9805351 A1

L2: Entry 3 of 3

File: EPAB

Feb 12, 1998

PUB-NO: WO009805351A1  
DOCUMENT-IDENTIFIER: WO 9805351 A1  
TITLE: METHODS FOR REGULATING GASTROINTESTINAL MOTILITY

PUBN-DATE: February 12, 1998

INVENTOR-INFORMATION:

NAME	COUNTRY
YOUNG, ANDREW A	US
GEDULIN, BRONISLAVA	US
BEELEY, NIGEL ROBERT ARNOLD	US
PRICKETT, KATHRYN S	US

INT-CL (IPC): A61 K 38/00; A61 K 38/26; G03 F 5/00; C07 K 2/00; C07 K 5/00  
EUR-CL (EPC): C07K014/575; A61K038/22

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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Term	Documents
GEDULIN-BRONISLAVA.DWPI,EPAB,USPT,PGPB.	3
GEDULIN-BRONISLAVAS	0
GEDULIN-BRONISLAVA.IN..USPT,PGPB,EPAB,DWPI,TDBD.	3
(GEDULIN-BRONISLAVA.IN.).USPT,PGPB,EPAB,DWPI,TDBD.	3

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Term	Documents
GEDULIN-BRONISLAVA.DWPI,EPAB,USPT,PGPB.	3
GEDULIN-BRONISLAVAS	0
GEDULIN-BRONISLAVA.IN..USPT,PGPB,EPAB,DWPI,TDBD.	3
(GEDULIN-BRONISLAVA.IN.).USPT,PGPB,EPAB,DWPI,TDBD.	3

Database:

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Search:

L2

[Refine Search](#)[Recall Text](#)[Clear](#)**Search History**DATE: Friday, January 24, 2003 [Printable Copy](#) [Create Case](#)Set Name  
side by sideQueryHit Count Set Name  
result set

DB=USPT,PGPB,EPAB,DWPI,TDBD; THES=ASSIGNEE; PLUR=YES;  
OP=ADJ

L2 gedulin-bronislava.in.3 L2L1 young-andrew-a\$.in.20 L1

END OF SEARCH HISTORY



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## Search Results - Record(s) 1 through 3 of 3 returned.

☐ 1. Document ID: US 20030004162 A1

L10: Entry 1 of 3

File: PGPB

Jan 2, 2003

PGPUB-DOCUMENT-NUMBER: 20030004162

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030004162 A1

TITLE: Use of glycogen phosphorylase inhibitors

PUBLICATION-DATE: January 2, 2003

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Treadway, Judith L.	Mystic	CT	US	

US-CL-CURRENT: 514/228.2; 514/233.5, 514/254.09, 514/365, 514/415

<a href="#">Full</a>	<a href="#">Title</a>	<a href="#">Citation</a>	<a href="#">Front</a>	<a href="#">Review</a>	<a href="#">Classification</a>	<a href="#">Date</a>	<a href="#">Reference</a>	<a href="#">Sequences</a>	<a href="#">Attachments</a>	<a href="#">Claims</a>	<a href="#">KWC</a>	<a href="#">Draw Desc</a>	<a href="#">Image</a>
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☐ 2. Document ID: US 20020187982 A1

L10: Entry 2 of 3

File: PGPB

Dec 12, 2002

PGPUB-DOCUMENT-NUMBER: 20020187982

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020187982 A1

TITLE: Glucagon antagonists/inverse agonists

PUBLICATION-DATE: December 12, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Behrens, Carsten	Kobenhavn N		DK	
Lau, Jesper	Farum		DK	
Madsen, Peter	Bagsvaerd		DK	

US-CL-CURRENT: 514/235.8; 514/254.03, 514/254.05, 514/326, 514/364, 514/381,  
514/567, 544/132, 544/138, 544/367, 546/208, 546/209, 548/132, 548/252, 562/442

<a href="#">Full</a>	<a href="#">Title</a>	<a href="#">Citation</a>	<a href="#">Front</a>	<a href="#">Review</a>	<a href="#">Classification</a>	<a href="#">Date</a>	<a href="#">Reference</a>	<a href="#">Sequences</a>	<a href="#">Attachments</a>	<a href="#">Claims</a>	<a href="#">KWC</a>	<a href="#">Draw Desc</a>	<a href="#">Image</a>
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☐ 3. Document ID: JP 2002538084 W WO 200041548 A2 AU 200024136 A NO 200103469  
A EP 1143989 A2 BR 200007823 A KR 2001086165 A KR 2002001719 A CN 1347327 A

L10: Entry 3 of 3

File: DWPI

Nov 12, 2002

DERWENT-ACC-NO: 2000-490999

DERWENT-WEEK: 200275

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TITLE: Lowering plasma glucagon using exendin, an exendin agonist, a modified exendin or a modified exendin agonist, useful for treating hyperglucagonemia and diabetes

INVENTOR: GEDULIN, B; YOUNG, A

PRIORITY-DATA: 2000US-175365P (January 10, 2000), 1999US-116380P (January 14, 1999), 1999US-132017P (April 30, 1999)

## PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
JP 2002538084 W	November 12, 2002		104	A61K038/00
WO 200041548 A2	July 20, 2000	E	096	A61K038/28
AU 200024136 A	August 1, 2000		000	A61K038/00
NO 200103469 A	September 14, 2001		000	A61K000/00
EP 1143989 A2	October 17, 2001	E	000	A61K038/00
BR 200007823 A	November 20, 2001		000	A61K038/00
KR 2001086165 A	September 8, 2001		000	A61K038/17
KR 2002001719 A	January 9, 2002		000	A61K038/22
CN 1347327 A	May 1, 2002		000	A61K038/22

INT-CL (IPC): A61 K 0/00; A61 K 38/00; A61 K 38/17; A61 K 38/22; A61 K 38/28; A61 K 45/00; A61 K 47/48; A61 P 5/00; A61 P 5/48; A61 P 17/00; A61 P 35/00; C07 K 14/435

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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Term	Documents
EXENDIN-4.DWPI,TDBD,EPAB,USPT,PGPB.	65
EXENDIN-4S	0
GLUCAGONOMA.DWPI,TDBD,EPAB,USPT,PGPB.	160
GLUCAGONOMAS.DWPI,TDBD,EPAB,USPT,PGPB.	63
(GLUCAGONOMA AND EXENDIN-4).USPT,PGPB,EPAB,DWPI,TDBD.	3
(EXENDIN-4 AND GLUCAGONOMA).USPT,PGPB,EPAB,DWPI,TDBD.	3

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L8: Entry 1 of 56

File: PGPB

Jan 16, 2003

PGPUB-DOCUMENT-NUMBER: 20030013646

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030013646 A1

TITLE: Methods to stimulate insulin production by pancreatic beta-cells

PUBLICATION-DATE: January 16, 2003

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Habener, Joel F.	Newton Centre	MA	US	
Thomas, Melissa K.	Boston	MA	US	

US-CL-CURRENT: 514/12; 424/93.21, 514/44[Full](#) [Title](#) [Citation](#) [Front](#) [Review](#) [Classification](#) [Date](#) [Reference](#) [Sequences](#) [Attachments](#)[KIMC](#) [Draw Desc](#) [Image](#)☐ 2. Document ID: US 20030008905 A1

L8: Entry 2 of 56

File: PGPB

Jan 9, 2003

PGPUB-DOCUMENT-NUMBER: 20030008905

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030008905 A1

TITLE: Method for the improvement of islet signaling in diabetes mellitus and for its prevention

PUBLICATION-DATE: January 9, 2003

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Demuth, Hans-Ulrich	Halle		DE	
Glund, Konrad	Halle		DE	

US-CL-CURRENT: 514/365; 514/423[Full](#) [Title](#) [Citation](#) [Front](#) [Review](#) [Classification](#) [Date](#) [Reference](#) [Sequences](#) [Attachments](#)[KIMC](#) [Draw Desc](#) [Image](#)☐ 3. Document ID: US 20030004162 A1

L8: Entry 3 of 56

File: PGPB

Jan 2, 2003

PGPUB-DOCUMENT-NUMBER: 20030004162

PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20030004162 A1

TITLE: Use of glycogen phosphorylase inhibitors

PUBLICATION-DATE: January 2, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Treadway, Judith L.	Mystic	CT	US	

US-CL-CURRENT: 514/228.2; 514/233.5, 514/254.09, 514/365, 514/415

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 4. Document ID: US 20020198242 A1

L8: Entry 4 of 56

File: PGPB

Dec 26, 2002

PGPUB-DOCUMENT-NUMBER: 20020198242  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20020198242 A1

TITLE: Method for the improvement of islet signaling in diabetes mellitus and for its prevention

PUBLICATION-DATE: December 26, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Demuth, Hans-Ulrich	Halle		DE	
Glund, Konrad	Halle		DE	

US-CL-CURRENT: 514/365; 514/423

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 5. Document ID: US 20020193390 A1

L8: Entry 5 of 56

File: PGPB

Dec 19, 2002

PGPUB-DOCUMENT-NUMBER: 20020193390  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20020193390 A1

TITLE: Pharmaceutical compositions containing an N-(substituted glycyI)-2-cyanopyrrolidine and at least one other antidiabetic agent and their use in inhibiting dipeptidyl peptidase-IV

PUBLICATION-DATE: December 19, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Villhauer, Edwin Bernard	Morristown	NJ	US	

US-CL-CURRENT: 514/275; 514/343, 514/423, 544/330, 546/279.1, 548/152, 548/537

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 6. Document ID: US 20020187982 A1

L8: Entry 6 of 56

File: PGPB

Dec 12, 2002

PGPUB-DOCUMENT-NUMBER: 20020187982

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020187982 A1

TITLE: Glucagon antagonists/inverse agonists

PUBLICATION-DATE: December 12, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Behrens, Carsten	Kobenhavn N		DK	
Lau, Jesper	Farum		DK	
Madsen, Peter	Bagsvaerd		DK	

US-CL-CURRENT: 514/235.8; 514/254.03, 514/254.05, 514/326, 514/364, 514/381,  
514/567, 544/132, 544/138, 544/367, 546/208 , 546/209, 548/132, 548/252, 562/442

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 7. Document ID: US 20020183369 A1

L8: Entry 7 of 56

File: PGPB

Dec 5, 2002

PGPUB-DOCUMENT-NUMBER: 20020183369

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020183369 A1

TITLE: Bicyclic pyrrolyl amides as glycogen phosphorylase inhibitors

PUBLICATION-DATE: December 5, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Du Bois, Daisy Joe	Palo Alto	CA	US	

US-CL-CURRENT: 514/367; 514/375, 514/393, 514/412, 548/153, 548/217, 548/303.1,  
548/453

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 8. Document ID: US 20020183367 A1

L8: Entry 8 of 56

File: PGPB

Dec 5, 2002

PGPUB-DOCUMENT-NUMBER: 20020183367

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020183367 A1

TITLE: 2,1-Oxazoline and 1,2-pyrazoline-based inhibitors of dipeptidyl peptidase IV

and method

PUBLICATION-DATE: December 5, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Sulsky, Richard B.	West Trenton	NJ	US	
Robl, Jeffrey A.	Newtown	PA	US	

US-CL-CURRENT: 514/365; 514/374, 514/385, 548/202, 548/215, 548/333.5

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 9. Document ID: US 20020164307 A1

L8: Entry 9 of 56

File: PGPB

Nov 7, 2002

PGPUB-DOCUMENT-NUMBER: 20020164307

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020164307 A1

TITLE: Stem cells of the islets of langerhans and their use in treating diabetes mellitus

PUBLICATION-DATE: November 7, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Habener, Joel F.	Newton Centre	MA	US	
Zulewski, Henryk	Basel	MA	CH	
Thomas, Melissa K.	Boston	MA	US	
Abraham, Elizabeth J.	Quincy	MA	US	
Vallejo, Mario	Madrid		ES	
Leech, Colin A.	Boston		US	

US-CL-CURRENT: 424/93.7; 424/93.21

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 10. Document ID: US 20020151065 A1

L8: Entry 10 of 56

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020151065

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020151065 A1

TITLE: Induction of beta cell differentiation in human cells by stimulation of the GLP-1 receptor

PUBLICATION-DATE: October 17, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Levine, Fred	Del Mar	CA	US	
Dufayet, Dominique	San Diego	CA	US	

US-CL-CURRENT: 435/455; 435/366

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 11. Document ID: US 20020146468 A1

L8: Entry 11 of 56

File: PGPB

Oct 10, 2002

PGPUB-DOCUMENT-NUMBER: 20020146468

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020146468 A1

TITLE: Extracts, compounds &amp; pharmaceutical compositions having anti-diabetic activity and their use

PUBLICATION-DATE: October 10, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Rubin, Ian Duncan	Nottingham	CT	GB	
Bindra, Jasjit Singh	Groton		US	
Cawthorne, Michael Anthony	Buckingham		GB	

US-CL-CURRENT: 424/725

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 12. Document ID: US 20020146405 A1

L8: Entry 12 of 56

File: PGPB

Oct 10, 2002

PGPUB-DOCUMENT-NUMBER: 20020146405

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020146405 A1

TITLE: Treatment of hibernating myocardium and diabetic cardiomyopathy with a GLP-1 peptide

PUBLICATION-DATE: October 10, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Coolidge, Thomas R.	Falls Village	CT	US	
Ehlers, Mario	Lincoln	NE	US	

US-CL-CURRENT: 424/94.61

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 13. Document ID: US 20020137666 A1

L8: Entry 13 of 56

File: PGPB

Sep 26, 2002

PGPUB-DOCUMENT-NUMBER: 20020137666

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020137666 A1

TITLE: USE OF EXENDINS AND AGONISTS THEREOF FOR THE REDUCTION OF FOOD INTAKE

PUBLICATION-DATE: September 26, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
BEELEY, NIGEL ROBERT ARNOLD	SOLANA BEACH	CA	US	
PRICKETT, KATHRYN S.	SAN DIEGO	CA	US	
BHAVSAR, SUNIL	SAN DIEGO	CA	US	

US-CL-CURRENT: 514/2; 514/12, 530/350

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 14. Document ID: US 20020123461 A1

L8: Entry 14 of 56

File: PGPB

Sep 5, 2002

PGPUB-DOCUMENT-NUMBER: 20020123461

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020123461 A1

TITLE: Enhancement of GLP-2 activity

PUBLICATION-DATE: September 5, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Drucker, Daniel J.	Toronto		CA	
Lovshin, Julie Ann Louise	Toronto		CA	

US-CL-CURRENT: 514/8; 514/12

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 15. Document ID: US 20020115605 A1

L8: Entry 15 of 56

File: PGPB

Aug 22, 2002

PGPUB-DOCUMENT-NUMBER: 20020115605

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020115605 A1

TITLE: Novel peptide with effects on cerebral health

PUBLICATION-DATE: August 22, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
During, Matthew	Philadelphia	PA	US	
Haile, Colin N.	Philadelphia	PA	US	

US-CL-CURRENT: 514/12; 514/16, 530/328



Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 16. Document ID: US 20020107206 A1

L8: Entry 16 of 56

File: PGPB

Aug 8, 2002

PGPUB-DOCUMENT-NUMBER: 20020107206

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020107206 A1

TITLE: Treatment of acute coronary syndrome with GLP-1

PUBLICATION-DATE: August 8, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Coolidge, Thomas R.	Falls Village	CT	US	
Ehlers, Mario	Lincoln	NE	US	

US-CL-CURRENT: 514/21

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 17. Document ID: US 20020099013 A1

L8: Entry 17 of 56

File: PGPB

Jul 25, 2002

PGPUB-DOCUMENT-NUMBER: 20020099013

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020099013 A1

TITLE: Active agent delivery systems and methods for protecting and administering active agents

PUBLICATION-DATE: July 25, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Piccariello, Thomas	Blacksburg	VA	US	
Olon, Lawrence P.	Bristol	TN	US	
Kirk, Randal J.	Radford	VA	US	

US-CL-CURRENT: 514/12

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 18. Document ID: US 20020098195 A1

L8: Entry 18 of 56

File: PGPB

Jul 25, 2002

PGPUB-DOCUMENT-NUMBER: 20020098195

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020098195 A1

TITLE: Effects of glucagon-like peptide-1 (7-36) on antro-pyloro-duodenal motility

PUBLICATION-DATE: July 25, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Goeke, Burkhard	Gauting		DE	
Schirra, Joerg	Kirchhain		DE	

US-CL-CURRENT: 424/184.1

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw. Desc	Image
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☐ 19. Document ID: US 20020065239 A1

L8: Entry 19 of 56

File: PGPB

May 30, 2002

PGPUB-DOCUMENT-NUMBER: 20020065239

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020065239 A1

TITLE: Methods and compositions for treatment of diabetes and related conditions via gene therapy

PUBLICATION-DATE: May 30, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Caplan, Shari L.	Sloatsburg	NY	US	
Boettcher, Brian R.	Morristown	NJ	US	
Slosberg, Eric D.	New York	NY	US	
Connelly, Sheila	Ijamsville	MD	US	
Kaleko, Michael	Rockville	MD	US	
Desai, Urvi J.	Germantown	MD	US	

US-CL-CURRENT: 514/44

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw. Desc	Image
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☐ 20. Document ID: US 20020058659 A1

L8: Entry 20 of 56

File: PGPB

May 16, 2002

PGPUB-DOCUMENT-NUMBER: 20020058659

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020058659 A1

TITLE: Imidazole compounds

PUBLICATION-DATE: May 16, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Andersen, Knud Erik	Brondby		DK	
Dorwald, Florencio Zaragiza	Ballerup		DK	
Peschke, Bernd	Malov		DK	
Sidemann, Ulla Grove	Valby		DK	
Rudolf, Klaus	Warthausen		DE	
Stenkamp, Dirk	Biberach		DE	
Hurnaus, Rudolf	Biberach		DE	
Muller, Stephan Georg	Warthausen		DE	
Krist, Bernd	Ulm		DE	
Eriksen, Birgitte	Farum		DE	

US-CL-CURRENT: 514/234.5; 514/314, 514/322, 514/338, 514/366, 514/394, 546/167,  
546/273.4, 548/159, 548/304.4

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 21. Document ID: US 20020049153 A1

L8: Entry 21 of 56

File: PGPB

Apr 25, 2002

PGPUB-DOCUMENT-NUMBER: 20020049153

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020049153 A1

TITLE: Long lasting insulinotropic peptides

PUBLICATION-DATE: April 25, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Bridon, Dominique P.	Outremont	CA	CA	
L'Archeveque, Benoit	Leval		CA	
Ezrin, Alan M.	Moraga		US	
Holmes, Darren L.	Montreal		CA	
Leblanc, Anouk	Montreal		CA	
St. Pierre, Serge	Ile Bizard		CA	

US-CL-CURRENT: 514/3; 514/12, 530/303

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 22. Document ID: US 20020037527 A1

L8: Entry 22 of 56

File: PGPB

Mar 28, 2002

PGPUB-DOCUMENT-NUMBER: 20020037527

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020037527 A1

TITLE: High density molecular arrays on porous surfaces

PUBLICATION-DATE: March 28, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Ellson, Richard N.	Palo Alto	CA	US	
Mutz, Mitchell W.	Palo Alto	CA	US	
Foote, James K.	Cupertino	CA	US	

US-CL-CURRENT: 435/6; 436/518

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 23. Document ID: US 20020037359 A1

L8: Entry 23 of 56

File: PGPB

Mar 28, 2002

PGPUB-DOCUMENT-NUMBER: 20020037359  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20020037359 A1

TITLE: Focused acoustic energy in the preparation of peptide arrays

PUBLICATION-DATE: March 28, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Mutz, Mitchell W.	Palo Alto	CA	US	
Ellson, Richard N.	Palo Alto	CA	US	

US-CL-CURRENT: 427/2.11; 435/176, 530/351, 530/388.1, 530/399

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 24. Document ID: US 20020019411 A1

L8: Entry 24 of 56

File: PGPB

Feb 14, 2002

PGPUB-DOCUMENT-NUMBER: 20020019411  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20020019411 A1

TITLE: Cyclopropyl-fused pyrrolidine-based inhibitors of dipeptidyl peptidase IV and method

PUBLICATION-DATE: February 14, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Robl, Jeffrey A.	Newtown	PA	US	
Sulsky, Richard B.	West Trenton	NJ	US	
Augeri, David J.	Princeton	NJ	US	
Magnin, David R.	Hamilton	NJ	US	
Hamann, Lawrence G.	Cherry Hill	NJ	US	
Betebenner, David A.	Lawrenceville	NJ	US	

US-CL-CURRENT: 514/299; 546/112

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 25. Document ID: US 20020010129 A1

L8: Entry 25 of 56

File: PGPB

Jan 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020010129  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20020010129 A1

TITLE: Shock heat treatment of polypeptides

PUBLICATION-DATE: January 24, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Matthiesen, Finn	Bronshoj		DK	

US-CL-CURRENT: 514/2; 530/350

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KMC	Draw Desc	Image
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☐ 26. Document ID: US 20010051646 A1

L8: Entry 26 of 56

File: PGPB

Dec 13, 2001

PGPUB-DOCUMENT-NUMBER: 20010051646  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010051646 A1

TITLE: Method for the improvement of islet signaling in diabetes mellitus and for its prevention

PUBLICATION-DATE: December 13, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Demuth, Hans-Ulrich	Halle		DE	
Glund, Konrad	Halle		DE	

US-CL-CURRENT: 514/369; 514/423

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KMC	Draw Desc	Image
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☐ 27. Document ID: US 20010049385 A1

L8: Entry 27 of 56

File: PGPB

Dec 6, 2001

PGPUB-DOCUMENT-NUMBER: 20010049385  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010049385 A1

TITLE: Imidazo heterocyclic compounds

PUBLICATION-DATE: December 6, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Andersen, Knud Erik	Brondby		DK	
Dorwald, Florencio Zaragoza	Ballerup		DK	
Peschke, Bernd	Malov		DK	

US-CL-CURRENT: 514/394; 548/303.1, 548/304.4, 548/304.7

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 28. Document ID: US 20010047084 A1

L8: Entry 28 of 56

File: PGPB

Nov 29, 2001

PGPUB-DOCUMENT-NUMBER: 20010047084  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010047084 A1

TITLE: Extendin derivatives

PUBLICATION-DATE: November 29, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Knudsen, Liselotte Bjerre	Valby		DK	
Huusfeldt, Per Olaf	Copenhagen K		DK	
Nielsen, Per Franklin	Vaerlose		DK	
Madsen, Kjeld	Vaerlose		DK	

US-CL-CURRENT: 530/399

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 29. Document ID: US 20010046489 A1

L8: Entry 29 of 56

File: PGPB

Nov 29, 2001

PGPUB-DOCUMENT-NUMBER: 20010046489  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010046489 A1

TITLE: Stem cells of the islets of langerhans and their use in treating diabetes mellitus

PUBLICATION-DATE: November 29, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Habener, Joel E.	Newton Center	MA	US	
Zulewski, Henryk	Geneva	MA	CH	
Abraham, Elizabeth J.	Quincy	MA	US	
Thomas, Melissa K.	Boston		US	
Vallejo, Mario	Madrid		ES	

US-CL-CURRENT: 424/93.21; 424/152.1, 435/366, 514/9

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 30. Document ID: US 20010038862 A1

L8: Entry 30 of 56

File: PGPB

Nov 8, 2001

PGPUB-DOCUMENT-NUMBER: 20010038862  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010038862 A1

TITLE: Topical and transdermal administration of peptidyl durgs using hydroxide releasing agents as permeation enhancers

PUBLICATION-DATE: November 8, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Luo, Eric C.	Plano	TX	US	
Hsu, Tsung-Min	San Diego	CA	US	

US-CL-CURRENT: 424/688; 514/2

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 31. Document ID: US 20010024824 A1

L8: Entry 31 of 56

File: PGPB

Sep 27, 2001

PGPUB-DOCUMENT-NUMBER: 20010024824  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010024824 A1

TITLE: Stem cells and their use in transplantation

PUBLICATION-DATE: September 27, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Moss, Peter Ian	London		GB	
Walters, David Martin	London		GB	
Pointer, Graham	London		GB	

US-CL-CURRENT: 435/366; 424/93.7

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 32. Document ID: US 20010012829 A1

L8: Entry 32 of 56

File: PGPB

Aug 9, 2001

PGPUB-DOCUMENT-NUMBER: 20010012829  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20010012829 A1

TITLE: Transepithelial delivery GLP-1 derivatives

PUBLICATION-DATE: August 9, 2001

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Anderson, Keith	San Diego	CA	US	
Agerso, Henrik	Fredensborg		DK	

US-CL-CURRENT: 514/12; 424/43

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KIMC	Draw Desc	Image
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☐ 33. Document ID: US 6506724 B1

L8: Entry 33 of 56

File: USPT

Jan 14, 2003

US-PAT-NO: 6506724

DOCUMENT-IDENTIFIER: US 6506724 B1

TITLE: Use of exendins and agonists thereof for the treatment of gestational diabetes mellitus

DATE-ISSUED: January 14, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Hiles; Richard A.	San Diego	CA		
Prickett; Kathryn S.	San Diego	CA		

US-CL-CURRENT: 514/2; 514/12, 514/3, 514/4, 514/866, 530/300, 530/324, 530/325

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KIMC	Draw Desc	Image
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☐ 34. Document ID: US 6500804 B2

L8: Entry 34 of 56

File: USPT

Dec 31, 2002

US-PAT-NO: 6500804

DOCUMENT-IDENTIFIER: US 6500804 B2

TITLE: Method for the improvement of islet signaling in diabetes mellitus and for its prevention

DATE-ISSUED: December 31, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Demuth; Hans-Ulrich	Halle			DE
Glund; Konrad	Halle			DE

US-CL-CURRENT: 514/19; 514/365, 514/866

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KIMC	Draw Desc	Image
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☐ 35. Document ID: US 6469021 B1

L8: Entry 35 of 56

File: USPT

Oct 22, 2002

US-PAT-NO: 6469021

DOCUMENT-IDENTIFIER: US 6469021 B1

TITLE: Non-peptide antagonists of GLP-1 receptor and methods of use

DATE-ISSUED: October 22, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Truesdale; Larry Kenneth	San Diego	CA		
Bychowski; Richard A.	Cardiff	CA		
Gonzalez; Javier	Oceanside	CA		
Kuki; Atsuo	Encinitas	CA		
Rajapakse; Ranjan Jagath	San Diego	CA		
Teng; Min	San Diego	CA		
Kiel; Dan	San Diego	CA		
Dhanoa; Daljit S.	West Chester	PA		
Hong; Yufeng	San Diego	CA		
Chou; Tso-sheng	San Diego	CA		
Ling; Anthony L.	San Diego	CA		
Johnson; Michael David	Cardiff	CA		
Gregor; Vlad Edward	San Diego	CA		

US-CL-CURRENT: 514/292; 514/232.8, 514/248, 514/287, 544/126, 544/233, 544/361,  
546/64, 546/85, 546/87

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 36. Document ID: US 6451974 B1

L8: Entry 36 of 56

File: USPT

Sep 17, 2002

US-PAT-NO: 6451974

DOCUMENT-IDENTIFIER: US 6451974 B1

TITLE: Method of acylating peptides and novel acylating agents

DATE-ISSUED: September 17, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Hansen; Louis Brammer	V.ae butt.ed.rl.o slashed.se			DK

US-CL-CURRENT: 530/345; 436/86, 436/90, 530/308, 530/333, 530/402

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 37. Document ID: US 6448045 B1

L8: Entry 37 of 56

File: USPT

Sep 10, 2002

US-PAT-NO: 6448045

DOCUMENT-IDENTIFIER: US 6448045 B1

TITLE: Inducing insulin gene expression in pancreas cells expressing recombinant PDX-1

DATE-ISSUED: September 10, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Levine; Fred	Del Mar	CA		
Dufayet; Dominique	San Diego	CA		

US-CL-CURRENT: 435/70.1; 435/455

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 38. Document ID: US 6437147 B1

L8: Entry 38 of 56

File: USPT

Aug 20, 2002

US-PAT-NO: 6437147

DOCUMENT-IDENTIFIER: US 6437147 B1

TITLE: Imidazole compounds

DATE-ISSUED: August 20, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Andersen; Knud Erik	Brondby			DK
Dorwald; Florencio Zaragoza	Ballerup			DK
Peschke; Bernd	Malov			DK
Sidelmann; Ulla Grove	Vedb.ae butted.k			DK
Rudolf; Klaus	Warthausen			DE
Stenkamp; Dirk	Birberach			DE
Hurnaus; Rudolf	Birberach			DE
Muller; Stephan Georg	Warthausen			DE
Krist; Bernd	Ulm			DE
Eriksen; Birgitte	Farum			DE

US-CL-CURRENT: 548/304.1; 548/302.7

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 39. Document ID: US 6432969 B1

L8: Entry 39 of 56

File: USPT

Aug 13, 2002

US-PAT-NO: 6432969

DOCUMENT-IDENTIFIER: US 6432969 B1

TITLE: N-(substituted glycy1)-2 cyanopyrrolidines, pharmaceutical compositions containing them and their use in inhibiting dipeptidyl peptidase-IV

DATE-ISSUED: August 13, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Villhauer; Edwin Bernard	Morristown	NJ		

US-CL-CURRENT: 514/275; 514/256, 514/343, 514/423, 544/332, 546/208, 546/279.1,  
548/540

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 40. Document ID: US 6399601 B1

L8: Entry 40 of 56

File: USPT

Jun 4, 2002

US-PAT-NO: 6399601

DOCUMENT-IDENTIFIER: US 6399601 B1

TITLE: Bicyclic pyrrolyl amides as glycogen phosphorylase inhibitors

DATE-ISSUED: June 4, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Du Bois; Daisy Joe	Palo Alto	CA		

US-CL-CURRENT: 514/233.8; 206/566, 514/321, 514/365, 514/419, 544/143, 546/198,  
548/181, 548/453

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 41. Document ID: US 6395767 B2

L8: Entry 41 of 56

File: USPT

May 28, 2002

US-PAT-NO: 6395767

DOCUMENT-IDENTIFIER: US 6395767 B2

TITLE: Cyclopropyl-fused pyrrolidine-based inhibitors of dipeptidyl peptidase IV and method

DATE-ISSUED: May 28, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Robl; Jeffrey A.	Newtown	PA		
Sulsky; Richard B.	West Trenton	NJ		
Augeri; David J.	Princeton	NJ		
Magnin; David R.	Hamilton	NJ		
Hamann; Lawrence G.	Cherry Hill	NJ		
Betebanner; David A.	Lawrenceville	NJ		

US-CL-CURRENT: 514/412; 548/452

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 42. Document ID: US 6376549 B1

L8: Entry 42 of 56

File: USPT

Apr 23, 2002

US-PAT-NO: 6376549

DOCUMENT-IDENTIFIER: US 6376549 B1

TITLE: Metformin-containing compositions for the treatment of diabetes

DATE-ISSUED: April 23, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Fine; Stuart A.	Northbrook	IL		
Kinsella; Kevin J.	La Jolla	CA		

US-CL-CURRENT: 514/635; 424/617, 424/626, 424/639, 424/655

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 43. Document ID: US 6329336 B1

L8: Entry 43 of 56

File: USPT

Dec 11, 2001

US-PAT-NO: 6329336

DOCUMENT-IDENTIFIER: US 6329336 B1

TITLE: Long lasting insulintropic peptides

DATE-ISSUED: December 11, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Bridon; Dominique P.	Outremont			CA
L'Archeveque; Benoit	Laval			CA
Ezrin; Alan M.	Moraga	CA		
Holmes; Darren L.	Montreal			CA
Leblanc; Anouk	Montreal			CA
St. Pierre; Serge	Ile Bizard			CA

US-CL-CURRENT: 514/2; 514/12

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 44. Document ID: US 6074875 A

L8: Entry 44 of 56

File: USPT

Jun 13, 2000

US-PAT-NO: 6074875

DOCUMENT-IDENTIFIER: US 6074875 A

TITLE: Materials and methods relating to the regulation of polypeptide production in cells

DATE-ISSUED: June 13, 2000

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Thorens; Bernard	Epalinge			CH

US-CL-CURRENT: 435/455; 435/325, 435/354, 435/358, 435/366

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KIMC	Draw Desc	Image
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☐ 45. Document ID: US 6051557 A

L8: Entry 45 of 56

File: USPT

Apr 18, 2000

US-PAT-NO: 6051557

DOCUMENT-IDENTIFIER: US 6051557 A

TITLE: Methods of enhancing functioning of the upper gastrointestinal tract

DATE-ISSUED: April 18, 2000

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Drucker; Daniel J.	Ontario			CA

US-CL-CURRENT: 514/12; 435/366, 530/308, 530/324

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KIMC	Draw Desc	Image
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☐ 46. Document ID: US 5846937 A

L8: Entry 46 of 56

File: USPT

Dec 8, 1998

US-PAT-NO: 5846937

DOCUMENT-IDENTIFIER: US 5846937 A

TITLE: Method of using exendin and GLP-1 to affect the central nervous system

DATE-ISSUED: December 8, 1998

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Drucker; Daniel J.	Toronto			CA

US-CL-CURRENT: 514/12; 514/2, 530/350, 530/399

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KIMC	Draw Desc	Image
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☐ 47. Document ID: US 5424286 A

L8: Entry 47 of 56

File: USPT

Jun 13, 1995

US-PAT-NO: 5424286

DOCUMENT-IDENTIFIER: US 5424286 A

TITLE: Exendin-3 and exendin-4 polypeptides, and pharmaceutical compositions comprising same

DATE-ISSUED: June 13, 1995

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Eng; John	Bronx	NY	10471	

US-CL-CURRENT: 514/2; 435/69.1, 514/866, 530/324

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 48. Document ID: DE 19921537 A1

L8: Entry 48 of 56

File: EPAB

Nov 23, 2000

PUB-NO: DE019921537A1

DOCUMENT-IDENTIFIER: DE 19921537 A1

TITLE: Treating carbohydrate metabolism disorders, especially diabetes, comprises activating insulin-secreting b-cells using glucagon-related peptide, glucose-dependent insulintropic polypeptide, exendin-4 or related drugs

PUBN-DATE: November 23, 2000

INVENTOR-INFORMATION:

NAME	COUNTRY
HOERSCH, DIETER	DE

INT-CL (IPC): A61 K 38/22; A61 K 38/26

EUR-CL (EPC): A61K038/26

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 49. Document ID: AU 200239384 A WO 200247716 A2

L8: Entry 49 of 56

File: DWPI

Jun 24, 2002

DERWENT-ACC-NO: 2002-519755

DERWENT-WEEK: 200267

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TITLE: Normalization of blood glucose levels, useful for e.g. treating hyperglycemia, diabetes, obesity, stroke or myocardial infarction, preventing cell deterioration or inducing weight loss, comprises use of glucagon-like peptides

INVENTOR: DODD, S W; MACE, K F ; TRAUTMANN, M E

PRIORITY-DATA: 2001US-298652P (June 15, 2001), 2000US-255251P (December 13, 2000), 2001US-295655P (June 4, 2001)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
AU 200239384 A	June 24, 2002		000	A61K038/26
WO 200247716 A2	June 20, 2002	E	087	A61K038/26

INT-CL (IPC): A61 K 38/26

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 50. Document ID: EP 1246638 A1 WO 200151078 A1 AU 200126380 A

L8: Entry 50 of 56

File: DWPI

Oct 9, 2002

DERWENT-ACC-NO: 2001-514422

DERWENT-WEEK: 200267

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TITLE: Use of exendin and exendin agonist compounds for modulating triglyceride levels, and treating heart disease and dyslipidemia

INVENTOR: KOLTERMAN, O G; YOUNG, A A

PRIORITY-DATA: 2000US-175365P (January 10, 2000)

## PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
EP 1246638 A1	October 9, 2002	E	000	A61K038/22
WO 200151078 A1	July 19, 2001	E	161	A61K038/22
AU 200126380 A	July 24, 2001		000	A61K038/22

INT-CL (IPC): A61 K 31/20; A61 K 31/22; A61 K 31/365; A61 K 31/40; A61 K 31:20; A61 K 31:22; A61 K 31:365; A61 K 31:40; A61 K 38/22; A61 K 38:22; A61 P 3/06; A61 K 38/22; A61 K 38/22; A61 K 38/22; A61 K 38/22; A61 K 31:40; A61 K 31:365; A61 K 31:22; A61 K 31:20

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 51. Document ID: EP 1257282 A1 WO 200139784 A1 AU 200118173 A US 20010024824 A1 US 20010046489 A1 US 20020164307 A1

L8: Entry 51 of 56

File: DWPI

Nov 20, 2002

DERWENT-ACC-NO: 2001-408256

DERWENT-WEEK: 200301

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TITLE: Treating diabetes mellitus or liver disease, comprises isolating a nestin-positive pancreatic stem cell from a pancreatic islet of a donor, and transferring the stem cell into the patient

INVENTOR: ABRAHAM, E J; FAUSTMAN, D ; HABENER, J L ; THOMAS, M K ; VALLEJO, M ; ZULEWSKI, H ; MOSS, P I ; POINTER, G ; WALTERS, D M ; HABENER, J E ; HABENER, J F ; LEECH, C A

PRIORITY-DATA: 2000US-238880P (October 6, 2000), 1999US-169082P (December 6, 1999), 2000US-215109P (June 28, 2000), 2000US-0731255 (December 6, 2000), 2000US-0731261 (December 6, 2000), 2001US-0963875 (September 26, 2001)

## PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
EP 1257282 A1	November 20, 2002	E	000	A61K035/00
WO 200139784 A1	June 7, 2001	E	102	A61K035/00
AU 200118173 A	June 12, 2001		000	A61K035/00
US 20010024824 A1	September 27, 2001		000	C12N005/08
US 20010046489 A1	November 29, 2001		000	A61K048/00
US 20020164307 A1	November 7, 2002		000	A61K048/00

INT-CL (IPC): A61 K 35/00; A61 K 39/395; A61 K 45/00; A61 K 48/00; C12 N 5/08; C12 N 15/85

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 52. Document ID: WO 200104156 A1 EP 1196444 A1 EP 1076066 A1 AU 200059660 A

L8: Entry 52 of 56

File: DWPI

Jan 18, 2001

DERWENT-ACC-NO: 2001-159381

DERWENT-WEEK: 200233

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TITLE: Novel peptide agonist of Glucagon-like peptide, useful for decreasing the level of blood glucose and for treating diseases like diabetes, obesity and eating disorders

INVENTOR: LARSEN, B D; MIKKELSEN, J D ; NEVE, S

PRIORITY-DATA: 1999EP-0610043 (August 9, 1999), 1999US-143591P (July 12, 1999)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
WO 200104156 A1	January 18, 2001	E	083	C07K014/575
EP 1196444 A1	April 17, 2002	E	000	C07K014/575
EP 1076066 A1	February 14, 2001	E	000	C07K014/575
AU 200059660 A	January 30, 2001		000	C07K014/575

INT-CL (IPC): A61 K 38/22; A61 K 47/48; A61 P 3/10; C07 K 14/575

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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☐ 53. Document ID: DE 19921537 A1

L8: Entry 53 of 56

File: DWPI

Nov 23, 2000

DERWENT-ACC-NO: 2001-050874

DERWENT-WEEK: 200107

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TITLE: Treating carbohydrate metabolism disorders, especially diabetes, comprises activating insulin-secreting b-cells using glucagon-related peptide, glucose-dependent insulinotropic polypeptide, exendin-4 or related drugs

INVENTOR: HOERSCH, D

PRIORITY-DATA: 1999DE-1021537 (May 11, 1999)

PATENT-FAMILY:



PUB-NO                      PUB-DATE                      LANGUAGE                      PAGES                      MAIN-IPC  
DE 19921537 A1              November 23, 2000                      010                      A61K038/22

INT-CL (IPC): A61 K 38/22; A61 K 38/26

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 54. Document ID: WO 200069911 A1 AU 200048555 A US 6329336 B1 EP 1180121 A1  
NO 200105584 A BR 200010750 A US 20020049153 A1 CN 1350548 A

L8: Entry 54 of 56

File: DWPI

Nov 23, 2000

DERWENT-ACC-NO: 2001-025008

DERWENT-WEEK: 200259

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TITLE: Novel modified insulintropic peptides for treating diabetes, nervous system disorders and for post surgery treatment, has reactive groups which react with amino, hydroxy or thiol groups on blood components

INVENTOR: BRIDON, D P; EZRIN, A M ; HOLMES, D L ; LARCHEVEQUE, B ; LEBLANC, A ; ST PIERRE, S ; L'ARCHEVEQUE, B ; BRIDON, D

PRIORITY-DATA: 1999US-159783P (October 15, 1999), 1999US-134406P (May 17, 1999), 2000US-0623618 (September 5, 2000), 2001US-0876388 (June 6, 2001)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
WO 200069911 A1	November 23, 2000	E	096	C07K014/605
AU 200048555 A	December 5, 2000		000	C07K014/605
US 6329336 B1	December 11, 2001		000	A01N037/18
EP 1180121 A1	February 20, 2002	E	000	C07K014/605
NO 200105584 A	January 3, 2002		000	C07K000/00
BR 200010750 A	February 26, 2002		000	C07K014/605
US 20020049153 A1	April 25, 2002		000	A61K038/28
CN 1350548 A	May 22, 2002		000	C07K014/605

INT-CL (IPC): A01 K 38/00; A01 N 37/18; A61 K 38/00; A61 K 38/26; A61 K 38/28; A61 P 3/08; C07 K 0/00; C07 K 14/575; C07 K 14/605; C07 K 14/62

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 55. Document ID: JP 2002538084 W WO 200041548 A2 AU 200024136 A NO  
200103469 A EP 1143989 A2 BR 200007823 A KR 2001086165 A KR 2002001719 A CN  
1347327 A

L8: Entry 55 of 56

File: DWPI

Nov 12, 2002

DERWENT-ACC-NO: 2000-490999

DERWENT-WEEK: 200275

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TITLE: Lowering plasma glucagon using exendin, an exendin agonist, a modified exendin or a modified exendin agonist, useful for treating hyperglucagonemia and diabetes

INVENTOR: GEDULIN, B; YOUNG, A

PRIORITY-DATA: 2000US-175365P (January 10, 2000), 1999US-116380P (January 14, 1999), 1999US-132017P (April 30, 1999)

## PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
JP 2002538084 W	November 12, 2002		104	A61K038/00
WO 200041548 A2	July 20, 2000	E	096	A61K038/28
AU 200024136 A	August 1, 2000		000	A61K038/00
NO 200103469 A	September 14, 2001		000	A61K000/00
EP 1143989 A2	October 17, 2001	E	000	A61K038/00
BR 200007823 A	November 20, 2001		000	A61K038/00
KR 2001086165 A	September 8, 2001		000	A61K038/17
KR 2002001719 A	January 9, 2002		000	A61K038/22
CN 1347327 A	May 1, 2002		000	A61K038/22

INT-CL (IPC): A61 K 0/00; A61 K 38/00; A61 K 38/17; A61 K 38/22; A61 K 38/28; A61 K 45/00; A61 K 47/48; A61 P 5/00; A61 P 5/48; A61 P 17/00; A61 P 35/00; C07 K 14/435

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 56. Document ID: US 5424286 A

L8: Entry 56 of 56

File: DWPI

Jun 13, 1995

DERWENT-ACC-NO: 1995-262627

DERWENT-WEEK: 199534

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TITLE: Stimulating/inhibiting insulin release with exendin polypeptide(s) - for treating diabetes mellitus and preventing hyperglycaemia.

INVENTOR: ENG, J

PRIORITY-DATA: 1993US-0066480 (May 24, 1993)

## PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
US 5424286 A	June 13, 1995		017	A61K038/16

INT-CL (IPC): A61 K 38/16; C07 K 14/46; C12 N 15/63

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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Generate Collection

Print

Term	Documents
EXENDIN-4.DWPI,TDBD,EPAB,USPT,PGPB.	65
EXENDIN-4S	0
GLUCAGON.DWPI,TDBD,EPAB,USPT,PGPB.	4444
GLUCAGONS.DWPI,TDBD,EPAB,USPT,PGPB.	49
(EXENDIN-4 AND GLUCAGON).USPT,PGPB,EPAB,DWPI,TDBD.	56
(EXENDIN-4 AND GLUCAGON).USPT,PGPB,EPAB,DWPI,TDBD.	56

**Display Format:****Change Format**[Previous Page](#)[Next Page](#)

## WEST Search History

DATE: Friday, January 24, 2003

**Set Name Query**

side by side

**Hit Count Set Name**

result set

*DB=USPT,PGPB,EPAB,DWPI,TDBD; THES=ASSIGNEE; PLUR=YES;  
OP=ADJ*

L10	exendin-4 and glucagonoma	3	L10
L9	exendin-4 and (necrolytic adj migratory adj erythema)	0	L9
L8	exendin-4 and glucagon	56	L8
L7	exendin-4 and ( glucagon and (necrolytic adj migratory adj erythema) and glucagonoma)	0	L7
L6	exendin-4	65	L6
L5	6348567.pn.	2	L5
L4	5348461.pn.	2	L4
L3	5846937.pn.	2	L3
L2	5424286.pn.	2	L2
L1	5424286.pn.	2	L1

END OF SEARCH HISTORY

**WEST**[Generate Collection](#)[Print](#)**Search Results - Record(s) 1 through 20 of 20 returned.**☐ 1. Document ID: US 20020141985 A1

L1: Entry 1 of 20

File: PGPB

Oct 3, 2002

PGPUB-DOCUMENT-NUMBER: 20020141985

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020141985 A1

TITLE: Peptide YY and peptide YY agonists for treatment of metabolic disorders

PUBLICATION-DATE: October 3, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Pittner, Richard A.	San Diego	CA	US	
<u>Young, Andrew A.</u>	La Jolla	CA	US	
Paterniti, James R. JR.	San Diego	CA	US	

US-CL-CURRENT: 424/94.1[Full](#) [Title](#) [Citation](#) [Front](#) [Review](#) [Classification](#) [Date](#) [Reference](#) [Sequences](#) [Attachments](#)[KMC](#) [Draw Desc](#) [Image](#)☐ 2. Document ID: US 20020010133 A1

L1: Entry 2 of 20

File: PGPB

Jan 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020010133

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020010133 A1

TITLE: METHOD FOR PREVENTING GASTRITIS USING AMYLIN OR AMYLIN AGONISTS

PUBLICATION-DATE: January 24, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
<u>YOUNG, ANDREW A.</u>	SAN DIEGO	CA	US	
GEDULIN, BRONISLAVA	SAN DIEGO	CA	US	
BEYNON, GARETH W.	BRIGHTWELL-CUM SOTWELL		UA	

US-CL-CURRENT: 514/12; 514/13, 514/14[Full](#) [Title](#) [Citation](#) [Front](#) [Review](#) [Classification](#) [Date](#) [Reference](#) [Sequences](#) [Attachments](#)[KMC](#) [Draw Desc](#) [Image](#)☐ 3. Document ID: US 6114304 A

L1: Entry 3 of 20

File: USPT

Sep 5, 2000

US-PAT-NO: 6114304

DOCUMENT-IDENTIFIER: US 6114304 A

TITLE: Methods for regulating gastrointestinal motility

DATE-ISSUED: September 5, 2000

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kolterman; Orville G.	Poway	CA		
<u>Young; Andrew A.</u>	Alpine	CA		
Rink; Timothy J.	La Jolla	CA		
Brown; Kathleen Ann Keiting	Wake Forest	NC		

US-CL-CURRENT: 514/12; 514/3

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 4. Document ID: US 6048514 A

L1: Entry 4 of 20

File: USPT

Apr 11, 2000

US-PAT-NO: 6048514

DOCUMENT-IDENTIFIER: US 6048514 A

TITLE: Amylin activity assays

DATE-ISSUED: April 11, 2000

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
<u>Young; Andrew A.</u>	San Diego	CA		
Cooper; Garth J. S.	Solana Beach	CA		
Rink; Timothy J.	La Jolla	CA		

US-CL-CURRENT: 424/9.2; 514/12, 514/21, 514/866, 514/884

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 5. Document ID: US 5814600 A

L1: Entry 5 of 20

File: USPT

Sep 29, 1998

US-PAT-NO: 5814600

DOCUMENT-IDENTIFIER: US 5814600 A

TITLE: Method and composition for treatment of insulin requiring mammals

DATE-ISSUED: September 29, 1998

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Rink; Timothy J.	La Jolla	CA		
<u>Young; Andrew A.</u>	Alpine	CA		

US-CL-CURRENT: 514/4; 514/12, 514/21, 514/3

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KMC	Draw Desc	Image
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☐ 6. Document ID: US 5739106 A

L1: Entry 6 of 20

File: USPT

Apr 14, 1998

US-PAT-NO: 5739106

DOCUMENT-IDENTIFIER: US 5739106 A

TITLE: Appetite regulating compositions

DATE-ISSUED: April 14, 1998

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Rink; Timothy J.	La Jolla	CA	92037	
<u>Young; Andrew A.</u>	Alpine	CA	91901	
Beeley; Nigel Robert Arnold	Solana Beach	CA	92037	
Prickett; Kathryn S.	San Diego	CA	92126	

US-CL-CURRENT: 514/12; 514/16, 514/18, 514/19, 530/303, 530/307, 530/312, 530/324, 530/328, 530/331

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KMC	Draw Desc	Image
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☐ 7. Document ID: US 5677279 A

L1: Entry 7 of 20

File: USPT

Oct 14, 1997

US-PAT-NO: 5677279

DOCUMENT-IDENTIFIER: US 5677279 A

TITLE: Methods and compositions for treating pain with amylin or agonists thereof

DATE-ISSUED: October 14, 1997

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
<u>Young; Andrew A.</u>	San Diego	CA		

US-CL-CURRENT: 514/12

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KMC	Draw Desc	Image
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☐ 8. Document ID: US 5656590 A

L1: Entry 8 of 20

File: USPT

Aug 12, 1997

US-PAT-NO: 5656590

DOCUMENT-IDENTIFIER: US 5656590 A

TITLE: Treatment of anorexia and related states

DATE-ISSUED: August 12, 1997

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Rink; Timothy J.	La Jolla	CA		
<u>Young; Andrew A.</u>	San Diego	CA		

US-CL-CURRENT: 514/3; 514/12, 514/4, 530/303

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 9. Document ID: US 5527771 A

L1: Entry 9 of 20

File: USPT

Jun 18, 1996

US-PAT-NO: 5527771

DOCUMENT-IDENTIFIER: US 5527771 A

TITLE: Methods and Compositions for treatment of diabetes mellitus, hypoglycemia &amp; other conditions

DATE-ISSUED: June 18, 1996

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Beaumont; Kevin	San Diego	CA		
<u>Young; Andrew A.</u>	San Diego	CA		

US-CL-CURRENT: 514/12; 530/307, 530/308

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 10. Document ID: US 5508260 A

L1: Entry 10 of 20

File: USPT

Apr 16, 1996

US-PAT-NO: 5508260

DOCUMENT-IDENTIFIER: US 5508260 A

TITLE: Methods and compositions for treatment of diabetes mellitus, hypoglycemia, and other conditions

DATE-ISSUED: April 16, 1996

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Beaumont; Kevin	San Diego	CA		
<u>Young; Andrew A.</u>	San Diego	CA		

US-CL-CURRENT: 514/4; 530/303, 530/307

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 11. Document ID: US 5376638 A

L1: Entry 11 of 20

File: USPT

Dec 27, 1994

US-PAT-NO: 5376638

DOCUMENT-IDENTIFIER: US 5376638 A

TITLE: Methods for treating renin-related disorders with amylin antagonists

DATE-ISSUED: December 27, 1994

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Young; Andrew A.	San Diego	CA		
Rink; Timothy J.	La Jolla	CA		

US-CL-CURRENT: 514/12; 514/11, 514/13

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 12. Document ID: US 5321008 A

L1: Entry 12 of 20

File: USPT

Jun 14, 1994

US-PAT-NO: 5321008

DOCUMENT-IDENTIFIER: US 5321008 A

TITLE: Methods and compositions for treatment of diabetes mellitus, hypoglycemia, and other conditions

DATE-ISSUED: June 14, 1994

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Beaumont; Kevin	San Diego	CA		
Young; Andrew A.	San Diego	CA		

US-CL-CURRENT: 514/4; 514/12, 514/21

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 13. Document ID: WO 9940788 A1

L1: Entry 13 of 20

File: EPAB

Aug 19, 1999

PUB-NO: WO009940788A1

DOCUMENT-IDENTIFIER: WO 9940788 A1

TITLE: INOTROPIC AND DIURETIC EFFECTS OF EXENDIN AND GLP-1

PUBN-DATE: August 19, 1999

## INVENTOR-INFORMATION:

NAME	COUNTRY
YOUNG, ANDREW A	US
VINE, WILL	US
BEELEY, NIGEL R A	US
PRICKETT, KATHRYN	US

INT-CL (IPC): A01 N 37/18  
EUR-CL (EPC): A61K031/00

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 14. Document ID: WO 9805351 A1

L1: Entry 14 of 20

File: EPAB

Feb 12, 1998

PUB-NO: WO009805351A1  
DOCUMENT-IDENTIFIER: WO 9805351 A1  
TITLE: METHODS FOR REGULATING GASTROINTESTINAL MOTILITY

PUBN-DATE: February 12, 1998

INVENTOR-INFORMATION:

NAME	COUNTRY
YOUNG, ANDREW A	US
GEDULIN, BRONISLAVA	US
BEELEY, NIGEL ROBERT ARNOLD	US
PRICKETT, KATHRYN S	US

INT-CL (IPC): A61 K 38/00; A61 K 38/26; G03 F 5/00; C07 K 2/00; C07 K 5/00  
EUR-CL (EPC): C07K014/575; A61K038/22

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 15. Document ID: WO 9640196 A1

L1: Entry 15 of 20

File: EPAB

Dec 19, 1996

PUB-NO: WO009640196A1  
DOCUMENT-IDENTIFIER: WO 9640196 A1  
TITLE: APPETITE REGULATING COMPOSITIONS

PUBN-DATE: December 19, 1996

INVENTOR-INFORMATION:

NAME	COUNTRY
RINK, TIMOTHY J	US
YOUNG, ANDREW A	US
BEELEY, NIGEL R A	US
PRICKETT, KATHERYN S	US

INT-CL (IPC): A61 K 38/00; C07 K 5/10; C07 K 7/06; C07 K 14/00  
EUR-CL (EPC): C07K014/575; A61K038/22, A61K038/22 , A61K038/23 , C07K014/595

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 16. Document ID: WO 9507098 A1

L1: Entry 16 of 20

File: EPAB

Mar 16, 1995

PUB-NO: WO009507098A1

DOCUMENT-IDENTIFIER: WO 9507098 A1

TITLE: METHODS FOR REGULATING GASTROINTESTINAL MOTILITY

PUBN-DATE: March 16, 1995

## INVENTOR-INFORMATION:

NAME

COUNTRY

KOLTERMAN, ORVILLE G

YOUNG, ANDREW A

RINK, TIMOTHY J

BROWN, KATHLEEN KEITING

INT-CL (IPC): A61 K 38/22; A61 K 38/23

EUR-CL (EPC): A61K038/22; A61K038/23

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Clip Img	Image
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☐ 17. Document ID: WO 9319774 A1

L1: Entry 17 of 20

File: EPAB

Oct 14, 1993

PUB-NO: WO009319774A1

DOCUMENT-IDENTIFIER: WO 9319774 A1

TITLE: AMYLIN AND POSSIBLY INSULIN CONTAINING COMPOSITION FOR THE TREATMENT OF ANOREXIA AND RELATED STATES

PUBN-DATE: October 14, 1993

## INVENTOR-INFORMATION:

NAME

COUNTRY

RINK, TIMOTHY J

YOUNG, ANDREW A

INT-CL (IPC): A61K 37/02

EUR-CL (EPC): A61K037/02; A61K038/22, A61K038/28

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 18. Document ID: WO 9220367 A1

L1: Entry 18 of 20

File: EPAB

Nov 26, 1992

PUB-NO: WO009220367A1

DOCUMENT-IDENTIFIER: WO 9220367 A1

TITLE: TREATMENT OF ANOREXIA AND RELATED STATES

PUBN-DATE: November 26, 1992

## INVENTOR-INFORMATION:

NAME	COUNTRY
RINK, TIMOTHY J	US
YOUNG, ANDREW A	US

INT-CL (IPC): A61K 37/02  
EUR-CL (EPC): A61K038/28; A61K038/22

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 19. Document ID: WO 9220366 A1

L1: Entry 19 of 20

File: EPAB

Nov 26, 1992

PUB-NO: WO009220366A1  
DOCUMENT-IDENTIFIER: WO 9220366 A1  
TITLE: INSULIN AND AMYLIN CONTAINING COMPOSITION FOR THE TREATMENT OF INSULIN DEFICIENT MAMMALS

PUBN-DATE: November 26, 1992

INVENTOR-INFORMATION:

NAME	COUNTRY
RINK, TIMOTHY J	US
YOUNG, ANDREW A	US

INT-CL (IPC): A61K 37/02  
EUR-CL (EPC): A61K037/02; A61K038/22, A61K038/28

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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☐ 20. Document ID: WO 9216222 A1

L1: Entry 20 of 20

File: EPAB

Oct 1, 1992

PUB-NO: WO009216222A1  
DOCUMENT-IDENTIFIER: WO 9216222 A1  
TITLE: METHODS AND COMPOSITIONS FOR TREATMENT OF DIABETES MELLITUS, HYPOGLYCEMIA, AND OTHER CONDITIONS

PUBN-DATE: October 1, 1992

INVENTOR-INFORMATION:

NAME	COUNTRY
BEAUMONT, KEVIN	US
YOUNG, ANDREW A	US

INT-CL (IPC): A61K 37/00; A61K 37/02; A61K 37/26; C07K 5/00; C07K 7/00; C07K 15/00; C07K 17/00  
EUR-CL (EPC): A61K038/28

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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KWIC	Draw Desc	Image
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(YOUNG-ANDREW-A\$.IN.).USPT,PGPB,EPAB,DWPI,TDBD.	20

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